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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification 6 : C07D 401/04, 401/14, 403/04, 403/12, 405/04, 409/12, 409/14, 487/04, A61K 31/44, 31/505 // (C07D 487/04, 239:00, 235:00) (C07D 487/04, 239:00, 239:00) (C07D 487/04, 243:00, 239:00)</p>		A3	<p>(11) International Publication Number: WO 98/24780 (43) International Publication Date: 11 June 1998 (11.06.98)</p>
<p>(21) International Application Number: PCT/US97/22949 (22) International Filing Date: 4 December 1997 (04.12.97)</p>		<p>(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</p>	
<p>(30) Priority Data: 60/032,128 5 December 1996 (05.12.96) US 60/050,950 13 June 1997 (13.06.97) US 08/976,053 21 November 1997 (21.11.97) US</p>		<p>Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p>	
<p>(71) Applicant (for all designated States except US): AMGEN INC. [US/US]; Amgen Center, 1840 De Havilland Drive, Thousand Oaks, CA 91320-1789 (US).</p>		<p>(88) Date of publication of the international search report: 30 July 1998 (30.07.98)</p>	
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<p>(74) Agents: ODRE, Steven, M. et al.; Amgen, Inc., Amgen Center, 1840 De Havilland Drive, Thousand Oaks, CA 91320-1789 (US).</p>			
<p>(54) Title: SUBSTITUTED PYRIMIDINONE AND PYRIDINONE COMPOUNDS AND THEIR USE</p>			
<p>(57) Abstract</p> <p>Selected novel substituted pyrimidinone and pyridone compounds are effective for prophylaxis and treatment of diseases, such as TNF-α, IL-1β, IL-6 and/or IL-8 mediated diseases, and other maladies, such as pain and diabetes. The invention encompasses novel compounds, analogs, prodrugs and pharmaceutically acceptable salts thereof, pharmaceutical compositions and methods for prophylaxis and treatment of diseases and other maladies or conditions involving inflammation, pain, diabetes and the like. The subject invention also relates to processes for making such compounds as well as to intermediates useful in such processes.</p>			

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INTERNATIONAL SEARCH REPORT

I. National Application No
PCT/US 97/22949

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 6 C07D401/04 C07D401/14 C07D403/04 C07D403/12 C07D405/04
 C07D409/12 C07D409/14 C07D487/04 A61K31/44 A61K31/505
 //((C07D487/04, 239:00, 235:00), (C07D487/04, 239:00, 239:00),

According to International Patent Classification(IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	C. A. MIRKIN ET AL.: JOURNAL OF THE AMERICAN CHEMICAL SOCIETY, vol. 112, no. 7, 1990, pages 2809-10, XP002064335 see page 2810, left-hand column, structure 3; right-hand column, table I, compound 3e ---	1-4
X	M. KOMATSU ET AL.: TETRAHEDRON LETTERS, vol. 22, no. 38, 1981, pages 3769-72, XP002064336 see page 3769, compound 6a ---	1-4
X	Y. OHSHIRO ET AL.: HETEROCYCLES, vol. 22, no. 3, 1984, pages 549-59, XP002064337 see page 551, compound 7 ---	1-4
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Further documents are listed in the continuation of box C.

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X document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

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1

Date of the actual completion of the international search	Date of mailing of the international search report
7 May 1998	02.06.98
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl. Fax: (+31-70) 340-3016	Authorized officer Hass, C

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 97/22949

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 (C07D487/04, 243:00, 239:00)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	P. I. MORTIMER: AUSTRALIAN JOURNAL OF CHEMISTRY, vol. 21, no. 2, 1968, pages 467-76, XP002064338 see page 468, compounds (VIII) and (X) ---	1-4
X	M. TAKAHASHI ET AL.: CHEMISTRY LETTERS, no. 6, 1987, pages 1229-32, XP002064339 see page 1231, compound 5 see page 1230, compound 8 ---	1-4
A	---	1-3, 20 -/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
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Date of the actual completion of the international search

7 May 1998

Date of mailing of the international search report

Name and mailing address of the ISA
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Authorized officer

Hass, C

INTERNATIONAL SEARCH REPORT

I. International Application No
PCT/US 97/22949

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	N. N. MAGDESIEVA ET AL.: CHEMISTRY OF HETEROCYCLIC COMPOUNDS, vol. 13, no. 9, 1978, pages 1177-80, XP002064340 (Translation from Khim. Geterotsikl. Soedin.) see page 1177, compounds IIIa - IIIe ----	1-4
X	R. D. YOUSSEFYEH ET AL.: JOURNAL OF THE CHEMICAL SOCIETY, PERKIN TRANSACTIONS 1, no. 23, 1974, pages 2649-54, XP002064341 see page 2651, compound (26) ----	1-4
X	DE 12 71 116 B (FARBENFABRIKEN BAYER AG) 17 June 1968 see columns 3-8, table ----	1-3, 12
X	J. J. BARR ET AL.: JOURNAL OF THE CHEMICAL SOCIETY, PERKIN TRANSACTIONS 1, no. 5, 1980, pages 1147-9, XP002064342 see page 1147, left-hand column, compound (5) ----	1-3, 12
X	M. TAKAHASHI ET AL.: HETEROCYCLES, vol. 22, no. 3, 1984, pages 581-4, XP002064343 see page 582, compounds 8a and 8e ----	1-3, 12
X	L. CAPUANO ET AL.: LIEBIGS ANNALEN DER CHEMIE, no. 4, 1991, pages 331-4, XP002064344 see page 331, compounds 8a, 8b, 8c ----	1-3, 12
X	T. L. GILCHRIST ET AL.: JOURNAL OF THE CHEMICAL SOCIETY, PERKIN TRANSACTIONS 1, no. 19, 1975, pages 1969-72, XP002064345 see page 1970, right-hand column, compound (11) ----	1-3, 12
X	T. L. GILCHRIST ET AL.: JOURNAL OF THE CHEMICAL SOCIETY, CHEMICAL COMMUNICATIONS, no. 12, 1974, pages 487-8, XP002064346 see page 488, compound (10) ----	1-3, 12
1		-/-

INTERNATIONAL SEARCH REPORT

I.	national Application No PCT/US 97/22949
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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	L. GIAMMANCO ET AL.: ATTI DELLA ACCADEMIA DI SCIENZE LETTERE E ARTI DI PALERMO, vol. 30, 1971, pages 93-107, XP002064347 see page 94, compound III; page 95, compounds VI, VIa, VIb, VIc, VId ---	1-3,12
X	H. YOSHIDA ET AL.: BULLETIN OF THE CHEMICAL SOCIETY OF JAPAN, vol. 56, no. 8, 1983, pages 2438-41, XP002064348 see page 2438, left-hand column, scheme 1, compounds 3 and 6; right-hand column, lines 6-14 ---	1-3,12
X	L. GIAMMANCO ET AL.: ANNALI DI CHIMICA, vol. 60, no. 3, 1970, pages 188-97, XP002064349 see page 190, table A; page 191, table B ---	1-3,12
X	L. GIAMMANCO: ATTI DELLA ACCADEMIA DI SCIENZE LETTERE E ARTI DI PALERMO, vol. 27, 1968, pages 469-83, XP002064350 see page 473, table A ---	1-3,12
A	WO 96 03387 A (G. D. SEARLE & CO.) 8 February 1996 see abstract; claims 1,6,10,15; examples ---	1,4,12, 28
A,P	WO 97 16442 A (MERCK & CO., INC.) 9 May 1997 cited in the application see abstract; claims 1,14,17-20,22,39 ---	1,4,12, 28
A,P	WO 97 12876 A (MERCK & CO., INC.) 10 April 1997 see claims 1,21,22,24,26 ---	1,4,12, 28
A	PATENT ABSTRACTS OF JAPAN vol. 10, no. 230 (C-365), 9 August 1996 & JP 61 063680 A (KANTO ISHI PHARMA CO LTD), 1 April 1986, see abstract ---	1,20,28
A	B. R. YREXA ET AL.: TETRAHEDRON, vol. 50, no. 21, 1994, pages 6173-80, XP002064351 see page 6174, compound 6; page 6176, compound 13; page 6178 ---	1-3,20
1		-/-

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 97/22949

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	E. KOTANI ET AL.: TETRAHEDRON, vol. 30, no. 17, 1974, pages 3027-30, XP002064352 see page 3027, compound 3 -----	1-3,20

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 97/22949

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
see FURTHER INFORMATION sheet PCT/ISA/210

2. Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210

3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.
 No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This international search report has not been established in respect of
-----the following reasons:

Claims Nos.: 29-46

because they relate to subject matter not required to be searched by this Authority, namely:

Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy

Claims Nos.: 1-26,28 (searched incompletely)

because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

The vast number of values for most of the variables, in conjunction with their cascading meanings and especially the presence of open definitions such as "substituted-aryl" or "heterocyclyl", render the scope of the invention for which protection is sought ill-defined and obscure. Consequently, an exhaustive and complete search is precluded for practical and economic reasons. The search was based upon though not limited to examples and tables given in the description (cf. Arts. 6, 15 and Rule 33 PCT).

Remark : Although claims 29-46 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/22949

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
DE 1271116 B			NONE
WO 9603387 A	08-02-96	US 5620999 A AU 3271695 A CA 2195846 A EP 0772601 A	15-04-97 22-02-96 08-02-96 14-05-97
WO 9716442 A	09-05-97	AU 1120897 A	22-05-97
WO 9712876 A	10-04-97	AU 7514396 A	28-04-97

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(74) Agents: ODRE, Steven, M. et al.; Amgen, Inc., Amgen Center, 1840 De Havilland Drive, Thousand Oaks, CA 91320-1789 (US).			
(54) Title: SUBSTITUTED PYRIMIDINONE AND PYRIDONE COMPOUNDS AND METHODS OF USE			
(57) Abstract			
<p>Selected novel substituted pyrimidinone and pyridone compounds are effective for prophylaxis and treatment of diseases, such as TNF-α, IL-1β, IL-6 and/or IL-8 mediated diseases, and other maladies, such as pain and diabetes. The invention encompasses novel compounds, analogs, prodrugs and pharmaceutically acceptable salts thereof, pharmaceutical compositions and methods for prophylaxis and treatment of diseases and other maladies or conditions involving inflammation, pain, diabetes and the like. The subject invention also relates to processes for making such compounds as well as to intermediates useful in such processes.</p>			

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EE	Estonia						

SUBSTITUTED PYRIMIDINONE AND PYRIDONE COMPOUNDS AND
METHODS OF USE

5

BACKGROUND OF THE INVENTION

This is a nonprovisional application derived from U.S. provisional application serial no. 60/032,128 filed December 5, 1996, U.S. provisional application serial no. 60/050,950 filed June 13, 1997 and U.S. 10 nonprovisional patent application serial no. not yet assigned filed November 21, 1997 each of which are incorporated herein by reference in their entirety. The present invention comprises a new class of compounds useful in treating diseases, such as TNF- α , IL-1 β , IL-6 15 and/or IL-8 mediated diseases and other maladies, such as pain and diabetes. In particular, the compounds of the invention are useful for the prophylaxis and treatment of diseases or conditions involving inflammation. This invention also relates to 20 intermediates and processes useful in the preparation of such compounds.

Interleukin-1 (IL-1) and Tumor Necrosis Factor α (TNF- α) are pro-inflammatory cytokines secreted by a variety of cells, including monocytes and macrophages, 25 in response to many inflammatory stimuli (e.g., lipopolysaccharide - LPS) or external cellular stress (e.g., osmotic shock and peroxide).

Elevated levels of TNF- α and/or IL-1 over basal levels have been implicated in mediating or exacerbating 30 a number of disease states including rheumatoid arthritis; Pagets disease; osteoporosis; multiple myeloma; uveitis; acute and chronic myelogenous leukemia; pancreatic β cell destruction; osteoarthritis; rheumatoid spondylitis; gouty arthritis; inflammatory 35 bowel disease; adult respiratory distress syndrome (ARDS); psoriasis; Crohn's disease; allergic rhinitis; ulcerative colitis; anaphylaxis; contact dermatitis;

asthma; muscle degeneration; cachexia; Reiter's syndrome; type I and type II diabetes; bone resorption diseases; graft vs. host reaction; ischemia reperfusion injury; atherosclerosis; brain trauma; multiple sclerosis; cerebral malaria; sepsis; septic shock; toxic shock syndrome; fever, and myalgias due to infection. HIV-1, HIV-2, HIV-3, cytomegalovirus (CMV), influenza, adenovirus, the herpes viruses (including HSV-1, HSV-2), and herpes zoster are also exacerbated by TNF- α .

It has been reported that TNF- α plays a role in head trauma, stroke, and ischemia. For instance, in animal models of head trauma (rat), TNF- α levels increased in the contused hemisphere (Shohami et al., *J. Cereb. Blood Flow Metab.* 14, 615 (1994)). In a rat model of ischemia wherein the middle cerebral artery was occluded, the levels of TNF- α mRNA of TNF- α increased (Feurstein et al., *Neurosci. Lett.* 164, 125 (1993)). Administration of TNF- α into the rat cortex has been reported to result in significant neutrophil accumulation in capillaries and adherence in small blood vessels. TNF- α promotes the infiltration of other cytokines (IL-1 β , IL-6) and also chemokines, which promote neutrophil infiltration into the infarct area (Feurstein, *Stroke* 25, 1481 (1994)). TNF- α has also been implicated to play a role in type II diabetes (Endocrinol. 130, 43-52, 1994; and Endocrinol. 136, 1474-1481, 1995).

TNF- α appears to play a role in promoting certain viral life cycles and disease states associated with them. For instance, TNF- α secreted by monocytes induced elevated levels of HIV expression in a chronically infected T cell clone (Clouse et al., *J. Immunol.* 142, 431 (1989)). Lahdevirta et al., (*Am. J. Med.* 85, 289 (1988)) discussed the role of TNF- α in the HIV associated states of cachexia and muscle degradation.

TNF- α is upstream in the cytokine cascade of inflammation. As a result, elevated levels of TNF- α may lead to elevated levels of other inflammatory and proinflammatory cytokines, such as IL-1, IL-6, and IL-8.

5 Elevated levels of IL-1 over basal levels have been implicated in mediating or exacerbating a number of disease states including rheumatoid arthritis; osteoarthritis; rheumatoid spondylitis; gouty arthritis; inflammatory bowel disease; adult respiratory distress
10 syndrome (ARDS); psoriasis; Crohn's disease; ulcerative colitis; anaphylaxis; muscle degeneration; cachexia; Reiter's syndrome; type I and type II diabetes; bone resorption diseases; ischemia reperfusion injury; atherosclerosis; brain trauma; multiple sclerosis;
15 sepsis; septic shock; and toxic shock syndrome. Viruses sensitive to TNF- α inhibition, e.g., HIV-1, HIV-2, HIV-3, are also affected by IL-1.

TNF- α and IL-1 appear to play a role in pancreatic β cell destruction and diabetes. Pancreatic β cells
20 produce insulin which helps mediate blood glucose homeostasis. Deterioration of pancreatic β cells often accompanies type I diabetes. Pancreatic β cell functional abnormalities may occur in patients with type II diabetes. Type II diabetes is characterized by a
25 functional resistance to insulin. Further, type II diabetes is also often accompanied by elevated levels of plasma glucagon and increased rates of hepatic glucose production. Glucagon is a regulatory hormone that attenuates liver gluconeogenesis inhibition by insulin.
30 Glucagon receptors have been found in the liver, kidney and adipose tissue. Thus glucagon antagonists are useful for attenuating plasma glucose levels (WO 97/16442, incorporated herein by reference in its entirety). By antagonizing the glucagon receptors, it
35 is thought that insulin responsiveness in the liver will

improve, thereby decreasing gluconeogenesis and lowering the rate of hepatic glucose production.

In rheumatoid arthritis models in animals, multiple intra-articular injections of IL-1 have led to an acute 5 and destructive form of arthritis (Chandrasekhar et al., *Clinical Immunol Immunopathol.* 55, 382 (1990)). In studies using cultured rheumatoid synovial cells, IL-1 is a more potent inducer of stromelysin than is TNF- α (Firestein, *Am. J. Pathol.* 140, 1309 (1992)). At sites 10 of local injection, neutrophil, lymphocyte, and monocyte emigration has been observed. The emigration is attributed to the induction of chemokines (e.g., IL-8), and the up-regulation of adhesion molecules (Dinarello, *Eur. Cytokine Netw.* 5, 517-531 (1994)).

15 IL-1 also appears to play a role in promoting certain viral life cycles. For example, cytokine-induced increase of HIV expression in a chronically infected macrophage line has been associated with a concomitant and selective increase in IL-1 production 20 (Folks et al., *J. Immunol.* 136, 40 (1986)). Beutler et al. (*J. Immunol.* 135, 3969 (1985)) discussed the role of IL-1 in cachexia. Baracos et al. (*New Eng. J. Med.* 308, 553 (1983)) discussed the role of IL-1 in muscle degeneration.

25 In rheumatoid arthritis, both IL-1 and TNF- α induce synoviocytes and chondrocytes to produce collagenase and neutral proteases, which leads to tissue destruction within the arthritic joints. In a model of arthritis (collagen-induced arthritis (CIA) in rats and mice), 30 intra-articular administration of TNF- α either prior to or after the induction of CIA led to an accelerated onset of arthritis and a more severe course of the disease (Brahn et al., *Lymphokine Cytokine Res.* 11, 253 (1992); and Cooper, *Clin. Exp. Immunol.* 898, 244 35 (1992)).

IL-8 has been implicated in exacerbating and/or causing many disease states in which massive neutrophil

infiltration into sites of inflammation or injury (e.g., ischemia) is mediated by the chemotactic nature of IL-8, including, but not limited to, the following: asthma, inflammatory bowel disease, psoriasis, adult respiratory distress syndrome, cardiac and renal reperfusion injury, thrombosis and glomerulonephritis. In addition to the chemotaxis effect on neutrophils, IL-8 also has the ability to activate neutrophils. Thus, reduction in IL-8 levels may lead to diminished neutrophil infiltration.

Several approaches have been taken to block the effect of TNF- α . One approach involves using soluble receptors for TNF- α (e.g., TNFR-55 or TNFR-75), which have demonstrated efficacy in animal models of TNF- α -mediated disease states. A second approach to neutralizing TNF- α using a monoclonal antibody specific to TNF- α , cA2, has demonstrated improvement in swollen joint count in a Phase II human trial of rheumatoid arthritis (Feldmann et al., *Immunological Reviews*, pp. 195-223 (1995)). These approaches block the effects of TNF- α and IL-1 by either protein sequestration or receptor antagonism.

US 5,100,897, incorporated herein by reference in its entirety, describes pyrimidinone compounds useful as angiotensin II antagonists wherein one of the pyrimidinone ring nitrogen atoms is substituted with a substituted phenylmethyl or phenethyl radical.

US 5,162,325, incorporated herein by reference in its entirety, describes pyrimidinone compounds useful as angiotensin II antagonists wherein one of the pyrimidinone ring nitrogen atoms is substituted with a substituted phenylmethyl radical.

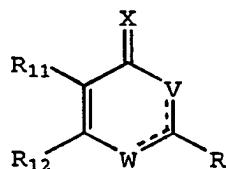
EP 481448, incorporated herein by reference in its entirety, describes pyrimidinone compounds useful as angiotensin II antagonists wherein one of the pyrimidinone ring nitrogen atoms is substituted with a substituted phenyl, phenylmethyl or phenethyl radical.

CA 2,020,370, incorporated herein by reference in its entirety, describes pyrimidinone compounds useful as angiotensin II antagonists wherein one of the pyrimidinone ring nitrogen atoms is substituted with a 5 substituted biphenylaliphatic hydrocarbon radical.

BRIEF DESCRIPTION OF THE INVENTION

The present invention comprises a new class of compounds useful in the prophylaxis and treatment of diseases, such as TNF- α , IL-1 β , IL-6 and/or IL-8 mediated diseases and other maladies, such as pain and diabetes. In particular, the compounds of the invention are useful for the prophylaxis and treatment of diseases or conditions involving inflammation. Accordingly, the 10 invention also comprises pharmaceutical compositions comprising the compounds, methods for the prophylaxis and treatment of TNF- α , IL-1 β , IL-6 and/or IL-8 mediated diseases, such as inflammatory, pain and diabetes 15 diseases, using the compounds and compositions of the invention, and intermediates and processes useful for the preparation of the compounds of the invention. 20

The compounds of the invention are represented by the following general structure:

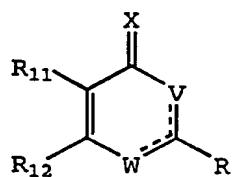


25 wherein the dashed lines represent a double bond between C(R) and V or W (i.e., -V=C(R)- or -W=C(R)-) and V, W, X, R, R¹¹ and R¹² are defined below.

The foregoing merely summarizes certain aspects of the invention and is not intended, nor should it be 30 construed, as limiting the invention in any way. All patents and other publications recited herein are hereby incorporated by reference in their entirety.

DETAILED DESCRIPTION OF THE INVENTION

In accordance with the present invention, there is provided compounds of the formula:

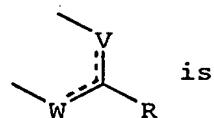


5 (I)

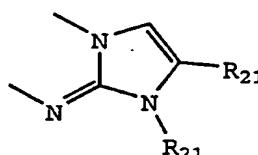
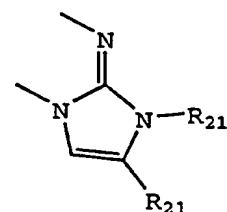
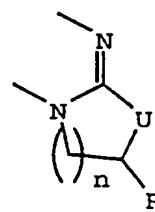
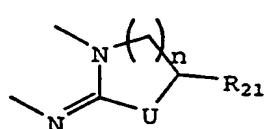
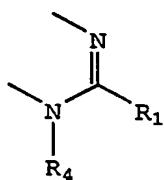
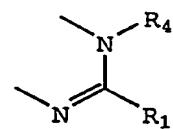
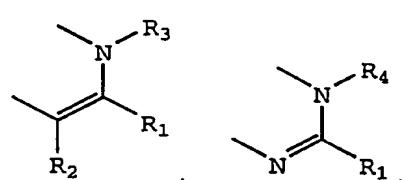
or a pharmaceutically acceptable salt thereof, wherein

X is O, S or NR₅; preferably, X is O or S; and most preferably, X is O;

10

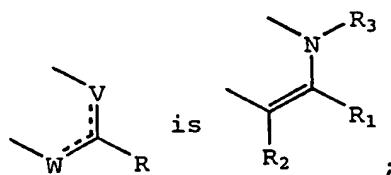


is

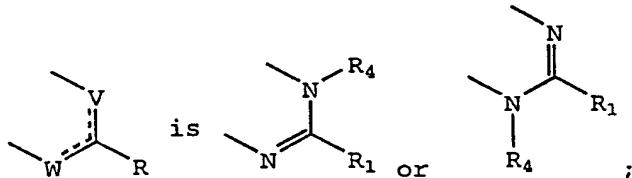


15 ; provided that the combined total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals in -VC(R)W- is 0-3, preferably, 0-2, most preferably, 0-1;

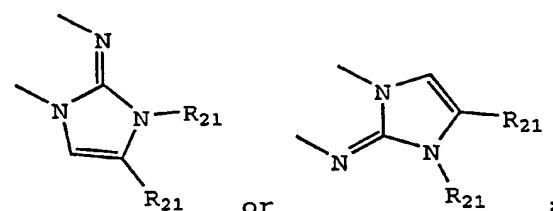
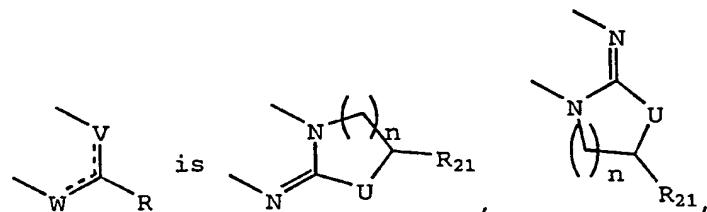
a first preferred subgroup of



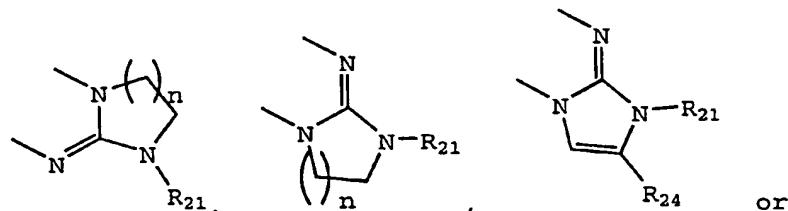
a second preferred subgroup of



5 a third preferred subgroup of



more preferably,

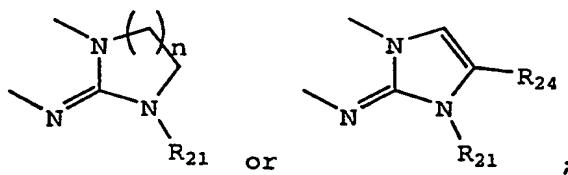


10

most preferably,



9



U is NR₂₁ or CHR₂₁; preferably, U is NR₂₁;

5 n is an integer of 1-3;

R₁ and R₂ are each independently -Y or -Z-Y, and R₃ and R₄ are each independently -Z-Y; provided that R₄ is other than a substituted-aryl, (substituted-aryl)methyl or (substituted-aryl)ethyl radical, and the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals in each -Y and -Z-Y is 0-3; preferably, 0-2; more preferably, 0-1;

15 preferably, R₂ is a radical of hydrogen, C₁-C₄ alkyl, halo, cyano, hydroxy, C₁-C₄ alkoxy, C₁-C₂ haloalkoxy of 1-3 halo radicals, C₁-C₄ alkylthio, amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino or C₁-C₂ haloalkyl of 1-3 halo radicals; more preferably, R₂ is a radical of hydrogen, C₁-C₄ alkyl, halo, cyano, hydroxy, C₁-C₄ alkoxy, trifluoromethoxy or trifluoromethyl; most preferably, R₂ is a hydrogen radical;

20 preferably, R₃ is a hydrogen radical or

25 (1) C₁-C₈ alkyl or C₂-C₈ alkenyl radical optionally substituted by (a) 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo, and (b)

30 1-2 radicals of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino,

10

(C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals; or
5 (2) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;

10 more preferably, R₃ is a hydrogen radical or
(1) C₁-C₈ alkyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy or aryl or heteroaryl
15 optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals; or
(2) aryl or heteroaryl radical optionally substituted by
20 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;
25 more preferably, R₃ is a hydrogen radical or C₁-C₈ alkyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy or aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;
30 more preferably, R₃ is a radical of hydrogen or C₁-C₄ alkyl; more preferably, R₃ is a hydrogen, methyl or ethyl radical;
35

preferably, R₄ is

(1) C₁-C₈ alkyl or C₂-C₈ alkenyl radical optionally substituted by (a) 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo, and (b) 1-2 radicals of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals; or
(2) heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;

more preferably, R₄ is
(1) C₁-C₈ alkyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy or aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals; or
(2) heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;

more preferably, R₄ is a C₁-C₈ alkyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy or aryl or heteroaryl optionally substituted by 1-3 radicals of
5 amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;

more preferably, R₄ is a C₁-C₄ alkyl radical; most
10 preferably, R₄ is a methyl or ethyl radical;

wherein each Z is independently a
(1) alkyl, alkenyl or alkynyl radical optionally substituted by (a) 1-3 radicals of amino, alkylamino,
15 dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio or halo, and (b) 1-2 radicals of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino,
20 alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, halo, alkyl or haloalkyl;
(2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, alkyl or haloalkyl; or
25 (3) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano, halo, alkyl or
30 haloalkyl;

preferably, each Z is independently a
(1) C₁-C₈ alkyl, C₂-C₈ alkenyl or C₂-C₈ alkynyl radical optionally substituted by (a) 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy) carbonylamino, C₁-C₄

alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo, and (b) 1-2 radicals of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄)alkylamino, C₁-C₅ alkanoylamino, (C₁-C₄)alkoxy carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; (2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄)alkylamino, C₁-C₅ alkanoylamino, (C₁-C₄)alkoxy carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or (3) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄)alkylamino, C₁-C₅ alkanoylamino, (C₁-C₄)alkoxy carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

more preferably, each Z is independently a (1) C₁-C₈ alkyl, C₂-C₈ alkenyl or C₂-C₈ alkynyl radical optionally substituted by (a) 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄)alkylamino, C₁-C₅ alkanoylamino, (C₁-C₄)alkoxy carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo, and (b) 1-2 radicals of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄)alkylamino, C₁-C₅ alkanoylamino, (C₁-C₄)alkoxy carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

(2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy,

5 C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or

(3) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy,

10 C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

more preferably, each Z is independently a

15 (1) C₁-C₈ alkyl or C₂-C₈ alkenyl radical optionally substituted by (a) 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo, and (b) 1-2 radicals of heterocyclyl,

20 aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3

25 halo radicals;

(2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, di-(C₁-C₄ alkyl)amino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl radicals; or

30 (3) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy,

35 C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

more preferably, each Z is independently a

(1) C₁-C₄ alkyl or C₂-C₅ alkenyl radical optionally

substituted by (a) 1-3 radicals of amino, di-(C₁-C₂

5 alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄

alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂

alkylthio or halo, and (b) 1-2 radicals of heterocyclyl,

aryl or heteroaryl optionally substituted by 1-3

radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₂

10 alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄

alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄

alkylthio, halo, C₁-C₄ alkyl or trifluoromethyl

radicals;

(2) heterocyclyl radical optionally substituted by 1-2

15 radicals of amino, di-(C₁-C₂ alkyl)amino, (C₁-C₄

alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂

alkylthio or C₁-C₄ alkyl radicals; or

(3) aryl or heteroaryl radical optionally substituted by

1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, C₁-C₅

20 alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-

C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or

trifluoromethyl radicals;

more preferably, each Z is independently a

25 (1) C₁-C₄ alkyl or C₂-C₅ alkenyl radical optionally

substituted by (a) 1-3 radicals of amino, di-(C₁-C₂

alkyl)amino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂

alkoxy, C₁-C₂ alkylthio or halo, and (b) 1-2 radicals of

aryl or heteroaryl optionally substituted by 1-2

30 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido,

(C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂

alkylthio, halo, C₁-C₄ alkyl or trifluoromethyl

radicals; or

(2) aryl or heteroaryl radical optionally substituted by

35 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido,

(C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂

alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

more preferably, each Z is independently a C₁-C₄ alkyl radical optionally substituted by 1-2 radicals of amino, di-(C₁-C₂ alkyl)amino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, halo or aryl or heteroaryl optionally substituted by 1-2 radicals of hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, halo, C₁-C₄ alkyl or trifluoromethyl radicals; and

most preferably, each Z is independently a C₁-C₄ alkyl radical optionally substituted by 1-2 radicals of amino, t-butoxycarbonylamino, dimethylamino, hydroxy, methoxy, methylthio or halo radicals;

each Y is independently a

- (1) hydrogen radical;
- (2) halo or nitro radical;
- 20 (3) -C(O)-R₂₀ or -C(NR₅)-NR₅R₂₁ radical;
- (4) -OR₂₁, -O-C(O)-R₂₁, -O-C(O)-NR₅R₂₁ or -O-C(O)-NR₂₂-S(O)₂-R₂₀ radical;
- (5) -SR₂₁, -S(O)-R₂₀, -S(O)₂-R₂₀, -S(O)₂-NR₅R₂₁, -S(O)₂-NR₂₂-C(O)-R₂₁, -S(O)₂-NR₂₂-C(O)-OR₂₀ or -S(O)₂-NR₂₂-C(O)-NR₅R₂₁ radical; or
- (6) -NR₅R₂₁, -NR₂₂-C(O)-R₂₁, -NR₂₂-C(O)-OR₂₀, -NR₂₂-C(O)-NR₅R₂₁, -NR₂₂-C(NR₅)-NR₅R₂₁, -NR₂₂-S(O)₂-R₂₀ or -NR₂₂-S(O)₂-NR₅R₂₁ radical;

30 preferably, each Y is independently a

- (1) hydrogen radical;
- (2) halo radical;
- (3) -C(O)-R₂₀ or -C(NR₅)-NR₅R₂₁ radical;
- (4) -OR₂₁, -O-C(O)-R₂₁ or -O-C(O)-NR₅R₂₁ radical;
- 35 (5) -SR₂₁, -S(O)-R₂₀, -S(O)₂-R₂₀ or -S(O)₂-NR₅R₂₁ radical; or

(6) $-NR_5R_{21}$, $-NR_{22}-C(O)-R_{21}$, $-NR_{22}-C(O)-OR_{20}$, $-NR_{22}-C(O)-NR_5R_{21}$, $-NR_{22}-C(NR_5)-NR_5R_{21}$, $-NR_{22}-S(O)_2-R_{20}$ or $-NR_{22}-S(O)_2-NR_5R_{21}$ radical;

5 more preferably, each Y is independently a

(1) hydrogen radical;

(2) $-C(O)-R_{20}$ radical;

(3) $-OR_{21}$, $-SR_{21}$, $-S(O)-R_{20}$, $-S(O)_2-R_{20}$ or $-S(O)_2-NR_5R_{21}$ radical; or

10 (4) $-NR_5R_{21}$, $-NR_{22}-C(O)-R_{21}$, $-NR_{22}-C(O)-OR_{20}$, $-NR_{22}-C(O)-NR_5R_{21}$, $-NR_{22}-S(O)_2-R_{20}$ or $-NR_{22}-S(O)_2-NR_5R_{21}$ radical;

more preferably, each Y is independently a

(1) hydrogen radical;

15 (2) $-C(O)-R_{20}$ radical;

(3) $-OR_{21}$, $-SR_{21}$, $-S(O)-R_{20}$, $-S(O)_2-R_{20}$ or $-S(O)_2-NR_5R_{21}$ radical; or

(4) $-NR_5R_{21}$, $-NR_{22}-C(O)-R_{21}$ or $-NR_{22}-S(O)_2-R_{20}$ radical;

20 more preferably, each Y is independently a

(1) $-C(O)-R_{20}$ radical;

(2) $-OR_{21}$, $-SR_{21}$, $-S(O)-R_{20}$, $-S(O)_2-R_{20}$ or $-S(O)_2-NR_5R_{21}$ radical; or

(3) $-NR_5R_{21}$, $-NR_{22}-C(O)-R_{21}$ or $-NR_{22}-S(O)_2-R_{20}$ radical.

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most preferably, each Y is independently a $-OR_{21}$, $-SR_{21}$ or $-NR_5R_{21}$ radical;

wherein each R₅ is independently

30 (1) hydrogen radicals;

(2) alkyl, alkenyl or alkynyl radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, hydroxy, alkoxy, alkylthio, $-SO_2H$ or halo; or

35 (3) aryl, heteroaryl, aralkyl, heteroaralkyl, heterocyclyl, heterocyclalkyl, cycloalkyl or

cycloalkylalkyl radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, hydroxy, alkoxy, alkylthio, alkyl or haloalkyl;

5 preferably, each R₅ is independently
(1) hydrogen radicals;
(2) C₁-C₈ alkyl, C₂-C₈ alkenyl or C₂-C₈ alkynyl radicals
optionally substituted by 1-3 radicals of amino, C₁-C₄
alkylamino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy,
10 C₁-C₄ alkylthio, -SO₂H or halo; or
(3) aryl, heteroaryl, aryl-C₁-C₄-alkyl, heteroaryl-C₁-C₄-
alkyl, heterocyclyl, heterocyclyl-C₁-C₄-alkyl, C₃-C₈
cycloalkyl or C₃-C₈-cycloalkyl-C₁-C₄-alkyl radicals
optionally substituted by 1-3 radicals of amino, C₁-C₄
15 alkylamino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy,
C₁-C₄ alkylthio, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3
halo radicals;

more preferably, each R₅ is independently
20 (1) hydrogen radicals;
(2) C₁-C₄ alkyl, C₂-C₅ alkenyl or C₂-C₅ alkynyl radicals
optionally substituted by 1-3 radicals of amino, C₁-C₄
alkylamino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy,
C₁-C₄ alkylthio, -SO₂H or halo; or
25 (3) aryl, heteroaryl, aryl-C₁-C₄-alkyl, heteroaryl-C₁-C₄-
alkyl, heterocyclyl, heterocyclyl-C₁-C₄-alkyl, C₃-C₈
cycloalkyl or C₃-C₈-cycloalkyl-C₁-C₄-alkyl radicals
optionally substituted by 1-3 radicals of amino, C₁-C₄
alkylamino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy,
30 C₁-C₄ alkylthio, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3
halo radicals;

more preferably, each R₅ is independently
(1) hydrogen radicals;

(2) C₁-C₄ alkyl or C₂-C₅ alkenyl radicals optionally substituted by 1-3 radicals of amino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, -SO₂H or halo; or

5 (3) phenyl-C₁-C₂-alkyl, heteroaryl-C₁-C₂-alkyl, heterocyclyl-C₁-C₂-alkyl or C₃-C₆-cycloalkyl-C₁-C₂-alkyl radicals optionally substituted by 1-3 radicals of amino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

more preferably, each R₅ is independently

(1) hydrogen radical;

(2) C₁-C₄ alkyl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂-alkyl)amino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio or halo; or

15 (3) phenyl-C₁-C₂-alkyl, heteroaryl-C₁-C₂-alkyl, heterocyclyl-C₁-C₂-alkyl or C₃-C₆-cycloalkyl-C₁-C₂-alkyl radicals optionally substituted by 1-3 radicals of amino, di-(C₁-C₂-alkyl)amino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, methoxy, methylthio, C₁-C₄ alkyl or trifluoromethyl radicals;

more preferably, each R₅ is independently

25 (1) hydrogen radical;

(2) C₁-C₄ alkyl radical optionally substituted by 1-3 halo radicals; or

(3) phenyl-C₁-C₂-alkyl or heteroaryl-C₁-C₂-alkyl, radicals optionally substituted by 1-3 radicals of

30 amino, dimethylamino, hydroxy, methoxy, methylthio, methyl or trifluoromethyl radicals;

more preferably, each R₅ is independently hydrogen or C₁-C₄ alkyl radical; and most preferably, each R₅ is a hydrogen radical;

wherein each R₂₀ is independently

(1) alkyl, alkenyl or alkynyl radicals optionally substituted by 1-3 radicals of amino, alkylamino,
5 dialkylamino, alkanoylamino, alkoxycarbonylamino, N-(alkoxycarbonyl)-N-(alkyl)amino, aminocarbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, halo or aralkoxy, aralkylthio, aralkylsulfonyl, cycloalkyl, heterocyclyl,
10 aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxycarbonylamino, alkylsulfonylamino, alkanoyl, hydroxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, halo, alkyl or haloalkyl;

15 (2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxycarbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, alkyl or haloalkyl; or
(3) aryl or heteroaryl radicals optionally substituted
20 by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxycarbonylamino, alkylsulfonylamino, alkoxycarbonyl, hydroxy, alkoxy, alkylthio, cyano, halo, azido, alkyl or haloalkyl;

25 preferably, each R₂₀ is independently

(1) C₁-C₈ alkyl, C₂-C₈ alkenyl or C₂-C₈ alkynyl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-
30 N-(C₁-C₄ alkyl)amino, aminocarbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo or aryl-C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-alkylsulfonyl, C₃-C₈ cycloalkyl, heterocyclyl, aryl or
35 heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄

alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, C₁-C₅ alkanoyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo, C₁-C₄ alkyl or 5 C₁-C₄ haloalkyl of 1-3 halo radicals;

(2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, 10 C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or

(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, 15 cyano, halo, azido, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

20 more preferably, each R₂₀ is independently

(1) C₁-C₈ alkyl, C₂-C₅ alkenyl or C₂-C₅ alkynyl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)- 25 N-(C₁-C₄ alkyl)amino, aminocarbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo or aryl-C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-alkylsulfonyl, C₃-C₈ cycloalkyl, heterocyclyl, aryl or 30 heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, C₁-C₅ alkanoyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄

alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;
(2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or
(3) aryl or heteroaryl radicals optionally substituted
by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

more preferably, each R₂₀ is independently
(1) C₁-C₈ alkyl or C₂-C₅ alkenyl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-C₄ alkyl)amino, aminocarbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo or aryl-C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-alkylsulfonyl, C₃-C₆ cycloalkyl, heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, C₁-C₅ alkanoyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;
(2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄

alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl; or
(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

more preferably, each R₂₀ is independently
(1) C₁-C₈ alkyl or C₂-C₅ alkenyl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-C₄ alkyl)amino, aminocarbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo or aryl-C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-alkylsulfonyl, C₃-C₆ cycloalkyl, heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, C₁-C₅ alkanoyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;
(2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, di-(C₁-C₄ alkyl)amino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl; or
(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, acetamido, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or trifluoromethyl radicals;

more preferably, each R₂₀ is independently
(1) C₁-C₈ alkyl radicals optionally substituted by 1-3
radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄
5 alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-C₄
alkyl)amino, aminocarbonylamino, hydroxy, C₁-C₄ alkoxy,
C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄
alkylsulfonyl, halo or C₃-C₆ cycloalkyl, heterocyclyl,
10 aryl or heteroaryl radicals optionally substituted by 1-
2 radicals of amino, di-(C₁-C₄ alkyl)amino, C₁-C₅
alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄
alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄
alkylthio, halo, C₁-C₄ alkyl or trifluoromethyl
15 radicals;
(2) heterocyclyl radical optionally substituted by 1-2
radicals of hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-
C₄ alkyl; or
(3) aryl or heteroaryl radicals optionally substituted
20 by 1-2 radicals of (C₁-C₄ alkoxy)carbonyl, amino, C₁-C₄
alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy,
C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or
trifluoromethyl radicals;
25 more preferably, each R₂₀ is independently
(1) C₁-C₆ alkyl radicals optionally substituted by 1-3
radicals of amino, methylamino, dimethylamino, t-
butoxycarbonylamino, N-((t-butoxy)carbonyl)-N-
(methyl)amino, aminocarbonylamino, hydroxy, butoxy,
30 methoxy, butylthio, methylthio, methylsulfinyl,
methylsulfonyl, halo or C₅-C₆ cycloalkyl, heterocyclyl,
phenyl or heteroaryl radicals optionally substituted by
1-2 radicals of amino, dimethylamino, acetamino,
hydroxy, methoxy, methylthio, halo, methyl or
35 trifluoromethyl radicals;

(2) heterocyclyl radical optionally substituted by 1-2 radicals of hydroxy or C₁-C₄ alkyl; or

(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, hydroxy,

5 methoxy, methylthio, halo, methyl or trifluoromethyl radicals;

more preferably, each R₂₀ is independently

(1) C₁-C₆ alkyl radicals optionally substituted by 1-3

10 radicals of amino, methylamino, dimethylamino, t-butoxycarbonylamino, N-((t-butoxy)carbonyl)-N-(methyl)amino, aminocarbonylamino, hydroxy, butoxy, methoxy, butylthio, methylthio, methylsulfinyl, methylsulfonyl, halo or C₅-C₆ cycloalkyl, heterocyclyl,

15 phenyl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, acetamino, hydroxy, methoxy, methylthio, halo, methyl or trifluoromethyl radicals;

(2) heterocyclyl radical; or

20 (3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, hydroxy, methoxy, methylthio, halo, methyl or trifluoromethyl radicals;

25 most preferably, each R₂₀ is independently

(1) C₁-C₆ alkyl radicals optionally substituted by 1-3 radicals of amino, methylamino, dimethylamino, hydroxy or phenyl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, hydroxy,

30 methoxy, methylthio, halo, methyl or trifluoromethyl radicals;

(2) heterocyclyl radical; or

(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, hydroxy,

35 methoxy, methylthio, halo, methyl or trifluoromethyl radicals;

each R₂₁ is independently hydrogen radical or R₂₀;

each R₂₂ is independently

(1) hydrogen radical;

5 (2) alkyl radical optionally substituted by a radical of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxycarbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, cyano, halo, alkyl or haloalkyl; or
10 (3) heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxycarbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio,

15 alkylsulfinyl, alkylsulfonyl, cyano, halo, alkyl or haloalkyl; provided when Z is a bond and Y is -NR₂₂-C(O)-NH₂, then R₂₂ is other then an optionally substituted aryl radical;

20 preferably, each R₂₂ is independently

(1) hydrogen radical;

(2) C₁-C₄ alkyl radical optionally substituted by a radical of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄) alkoxy carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or

25 (3) heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄) alkoxy carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; provided when Z is a

bond and Y is $-NR_{22}-C(O)-NH_2$, then R₂₂ is other than an optionally substituted aryl radical;

more preferably, each R₂₂ is independently

- 5 (1) hydrogen radical; or
- (2) C₁-C₄ alkyl radical optionally substituted by a radical of phenyl or heteroaryl optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

more preferably, each R₂₂ is independently hydrogen or C₁-C₄ alkyl radical; and most preferably, each R₂₂ is

- 15 independently hydrogen or methyl radical;

R₁₁ and R₁₂ are each independently an aryl or heteroaryl radical optionally substituted by 1-3 radicals of

- (1) R₃₀;
- 20 (2) halo or cyano radicals;
- (3) $-C(O)-R_{30}$, $-C(O)-OR_{29}$, $-C(O)-NR_{31}R_{32}$ or $-C(NR_{31})-$ NR₃₁R₃₂ radicals;
- (4) $-OR_{29}$, $-O-C(O)-R_{29}$, $-O-C(O)-NR_{31}R_{32}$ or $-O-C(O)-NR_{33}-$ S(O)₂-R₃₀ radicals;
- 25 (5) $-SR_{29}$, $-S(O)-R_{30}$, $-S(O)_2-R_{30}$, $-S(O)_2-NR_{31}R_{32}$, $-S(O)_2-$ NR₃₃-C(O)-R₃₀, $-S(O)_2-NR_{33}-C(O)-OR_{30}$ or $-S(O)_2-NR_{33}-C(O)-$ NR₃₁R₃₂ radicals; or
- (6) $-NR_{31}R_{32}$, $-NR_{33}-C(O)-R_{29}$, $-NR_{33}-C(O)-OR_{30}$, $-NR_{33}-C(O)-$ NR₃₁R₃₂, $-NR_{33}-C(NR_{31})-NR_{31}R_{32}$, $-NR_{33}-S(O)_2-R_{30}$ or $-NR_{33}-$ S(O)₂-NR₃₁R₃₂ radicals;
- 30 provided that (1) R₁₁ is other than a 4-pyridyl, 4-pyrimidinyl, 4-quinolyl or 6-isoquinolinalyl radical optionally substituted by 1-2 substituents; and (2) the total number of aryl, heteroaryl, cycloalkyl and heterocyclal radicals substituted on each of R₁₁ and R₁₂ is 0-1;

preferably, R₁₁ and R₁₂ are each independently an aryl or heteroaryl radical optionally substituted by 1-2 radicals of

- 5 (1) R₃₀;
- (2) halo or cyano radicals;
- (3) -C(O)-R₃₀, -C(O)-OR₂₉, -C(O)-NR₃₁R₃₂ or -C(NR₃₁)-NR₃₁R₃₂ radicals;
- (4) -OR₂₉, -O-C(O)-R₂₉, -O-C(O)-NR₃₁R₃₂ or -O-C(O)-NR₃₃-

10 S(O)₂-R₃₀ radicals;

- (5) -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -S(O)₂-NR₃₃-C(O)-R₃₀, -S(O)₂-NR₃₃-C(O)-OR₃₀ or -S(O)₂-NR₃₃-C(O)-
- 15 NR₃₁R₃₂ radicals; or

- (6) -NR₃₁R₃₂, -NR₃₃-C(O)-R₂₉, -NR₃₃-C(O)-OR₃₀, -NR₃₃-C(O)-
- 15 NR₃₁R₃₂, -NR₃₃-C(NR₃₁)-NR₃₁R₃₂, -NR₃₃-S(O)₂-R₃₀ or -NR₃₃-
- S(O)₂-NR₃₁R₃₂ radicals;

provided that (1) R₁₁ is other than a 4-pyridyl, 4-pyrimidinyl, 4-quinolyl or 6-isouquinolinyl radical optionally substituted by 1-2 substituents; and (2) the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each of R₁₁ and R₁₂ is 0-1;

more preferably, R₁₁ and R₁₂ are each independently an aryl or heteroaryl radical optionally substituted by 1-2 radicals of

- (1) R₃₀;
- (2) halo or cyano radicals;
- (3) -C(O)-R₃₀, -C(O)-OR₂₉, -C(O)-NR₃₁R₃₂ or -C(NR₃₁)-NR₃₁R₃₂ radicals; or
- (4) -OR₂₉, -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -NR₃₁R₃₂, -NR₃₃-C(O)-R₂₉ or -NR₃₃-C(O)-OR₃₀ radicals;

more preferably, R₁₁ is an aryl radical and R₁₂ is a heteroaryl radical, wherein the aryl and heteroaryl radicals are optionally substituted by 1-2 radicals of

(1) R_{30} ;
(2) halo or cyano radicals;
(3) $-C(O)-R_{30}$, $-C(O)-OR_{29}$, $-C(O)-NR_{31}R_{32}$ or $-C(NR_{31})-$
NR₃₁R₃₂ radicals; or
5 (4) $-OR_{29}$, $-SR_{29}$, $-S(O)-R_{30}$, $-S(O)_2-R_{30}$, $-S(O)_2-NR_{31}R_{32}$,
 $-NR_{31}R_{32}$ or $-NR_{33}-C(O)-R_{29}$ radicals;

more preferably, R_{11} is an aryl radical and R_{12} is a heteroaryl radical, wherein the aryl and heteroaryl
10 radicals are optionally substituted by 1-2 radicals of

(1) R_{30} ;
(2) halo or cyano radicals; or
(3) $-C(O)-NR_{31}R_{32}$, $-OR_{29}$, $-SR_{29}$, $-S(O)-R_{30}$, $-S(O)_2-R_{30}$, $-S(O)_2-NR_{31}R_{32}$,
15 $-NR_{31}R_{32}$ or $-NR_{33}-C(O)-R_{29}$ radicals;

more preferably, R_{11} is an aryl radical optionally substituted by 1-2 radicals of (1) R_{30} ; (2) halo or cyano radicals; or (3) $-C(O)-NR_{31}R_{32}$, $-OR_{29}$, $-SR_{29}$, $-S(O)-R_{30}$, $-S(O)_2-R_{30}$, $-S(O)_2-NR_{31}R_{32}$, $-NR_{31}R_{32}$ or $-NR_{33}-C(O)-R_{29}$ radicals; more preferably, R_{11} is an aryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methylthio, methylsulfinyl, methylsulfonyl, aminocarbonyl, methyl or trifluoromethyl radicals; more
20 preferably, R_{11} is an unsubstituted phenyl or naphthyl radical or a phenyl radical substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methylthio, methylsulfinyl, methylsulfonyl, aminocarbonyl, methyl or trifluoromethyl radicals; more
25 preferably, R_{11} is an unsubstituted phenyl or naphthyl radical or a phenyl radical substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methylthio, methylsulfinyl, methylsulfonyl, aminocarbonyl, methyl or trifluoromethyl radicals; and most preferably, R_{11} is an unsubstituted phenyl radical or a phenyl radical substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methylthio, methylsulfinyl, methylsulfonyl, methyl or trifluoromethyl radicals;

more preferably, R₁₂ is a heteroaryl radical optionally substituted by 1-2 radicals of (1) R₃₀; (2) halo or cyano radicals; or (3) -C(O)-NR₃₁R₃₂, -OR₂₉, -SR₂₉, -NR₃₁R₃₂ or -NR₃₃-C(O)-R₂₉ radicals; more preferably, R₁₂ is a heteroaryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methyl or trifluoromethyl radicals; more preferably, R₁₂ is a 4-pyridyl, 4-quinolinyl, 4-imidazolyl or 4-pyrimidinyl radical 5 optionally substituted by a radical of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methyl or trifluoromethyl radicals; and most preferably, R₁₂ is a 4-pyridyl radical optionally substituted by a radical of amino, dimethylamino, acetamido, hydroxy, 10 halo, cyano, methoxy, methyl or trifluoromethyl radicals; R₁₂ is a 4-pyridyl radical optionally substituted by a radical of amino, dimethylamino, acetamido, hydroxy, 15 halo, cyano, methoxy, methyl or trifluoromethyl radicals;

wherein each R₃₀ is independently
(1) alkyl, alkenyl or alkynyl radicals optionally 20 substituted by 1-3 radicals of -NR₃₁R₃₁, -CO₂R₂₃, hydroxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, cyano, halo or aralkoxy, aralkylthio, aralkylsulfonyl, heterocyclyl, aryl or heteroaryl 25 radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, cyano, halo, alkyl or haloalkyl;
(2) heterocyclyl radical optionally substituted by 1-3 30 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano, alkyl or haloalkyl; or
(3) aryl or heteroaryl radicals optionally substituted 35 by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino,

hydroxy, alkoxy, alkylthio, cyano, halo, alkyl or haloalkyl;

preferably, each R₃₀ is independently

- 5 (1) C₁-C₄ alkyl, C₂-C₄ alkenyl or C₂-C₄ alkynyl radicals optionally substituted by 1-3 radicals of -NR₃₁R₃₁, -CO₂R₂₃, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, cyano, halo or aryl-C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-alkylsulfonyl, heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;
- 10 (2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or
- 15 (3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;
- 20 more preferably, each R₃₀ is independently
(1) C₁-C₄ alkyl radical optionally substituted by 1-3 radicals of
(a) -NR₃₁R₃₁;
- 25 (b) C₁-C₄ alkoxy-carbonyl or phenoxy carbonyl or
30 phenylmethoxycarbonyl optionally substituted by 1-3

radicals of amino, alkylamino, di-(C₁-C₄-alkyl)amino,
C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄
alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄
alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl;

5 or

(c) hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, or phenyl-C₁-
C₄-alkoxy, phenyl-C₁-C₄-alkylthio, heterocyclyl, phenyl
or heteroaryl radicals optionally substituted by 1-3
radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄
alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄
alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of
1-3 halo radicals;

(2) C₁-C₄ haloalkyl of 1-3 halo radical; or

15 (3) aryl or heteroaryl radicals optionally substituted
by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄
alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄
alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl

20 radicals;

more preferably, each R₃₀ is independently

(1) C₁-C₄ alkyl radical optionally substituted by

(a) amino, C₁-C₄ alkylamino or di-(C₁-C₄-alkyl)amino
25 radicals; or

(b) hydroxy, C₁-C₄ alkoxy, heterocyclyl, phenyl or
heteroaryl radicals optionally substituted by 1-3
radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄
alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄
alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl
30 radicals;

(2) C₁-C₂ haloalkyl of 1-3 halo radical; or

(3) aryl or heteroaryl radicals optionally substituted
35 by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄

alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

5

more preferably, each R₃₀ is independently
(1) C₁-C₄ alkyl radical optionally substituted by a phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, hydroxy, C₁-C₂ alkoxy, halo, C₁-C₄ alkyl or trifluoromethyl radicals;
(2) trifluoromethyl radical; or
(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, hydroxy, C₁-C₂ alkoxy, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

more preferably, each R₃₀ is independently
(1) C₁-C₄ alkyl radical optionally substituted by a phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;
(2) trifluoromethyl radical; or
(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;

most preferably, R₃₀ is independently
(1) C₁-C₄ alkyl radical optionally substituted by a phenyl or heteroaryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;
(2) trifluoromethyl radical; or

(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;

5

each R₂₉ is independently hydrogen radical or R₃₀; and most preferably, R₂₉ is an aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;

10

each R₃₁ is independently

(1) hydrogen radicals;

(2) alkyl radical optionally substituted by an

15 cycloalkyl, aryl, heterocyclyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano, alkyl or haloalkyl; or

20 (3) aryl, heteroaryl, heterocyclyl or cycloalkyl radical optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano, alkyl or haloalkyl;

25

preferably, each R₃₁ is independently

(1) hydrogen radicals;

(2) C₁-C₄ alkyl radical optionally substituted by an C₃-C₈ cycloalkyl, aryl, heterocyclyl or heteroaryl radical

30 optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or

35 (3) aryl, heteroaryl, heterocyclyl or C₃-C₈ cycloalkyl radical optionally substituted by 1-3 radicals of amino,

C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3
5 halo radicals;

more preferably, each R₃₁ is independently
(1) hydrogen radicals; or
(2) C₁-C₄ alkyl radical optionally substituted by an
10 phenyl or heteroaryl radical optionally substituted by
1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or trifluoromethyl
15 radicals;

more preferably, each R₃₁ is independently hydrogen or
C₁-C₄ alkyl radicals; and most preferably, each R₃₁ is
independently hydrogen, methyl or ethyl radicals;

20 each R₃₂ is independently
(1) hydrogen radicals;
(2) alkyl radical optionally substituted by an
cycloalkyl, aryl, heterocyclyl or heteroaryl radical
25 optionally substituted by 1-3 radicals of amino,
alkylamino, dialkylamino, alkanoylamino,
alkoxycarbonylamino, alkylsulfonylamino, hydroxy,
alkoxy, alkylthio, cyano, alkyl or haloalkyl; or
(3) aryl, heteroaryl, heterocyclyl or cycloalkyl radical
30 optionally substituted by 1-3 radicals of amino,
alkylamino, dialkylamino, alkanoylamino,
alkoxycarbonylamino, alkylsulfonylamino, hydroxy,
alkoxy, alkylthio, cyano, alkyl or haloalkyl;

35 preferably, each R₃₂ is independently
(1) hydrogen radicals;

(2) C₁-C₄ alkyl radical optionally substituted by an C₃-C₈ cycloalkyl, aryl, heterocyclyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino,
5 (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or
(3) aryl, heteroaryl, heterocyclyl or C₃-C₈ cycloalkyl radical optionally substituted by 1-3 radicals of amino,
10 C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;
15 more preferably, each R₃₂ is independently
(1) hydrogen radicals;
(2) C₁-C₄ alkyl radical optionally substituted by an C₃-C₆ cycloalkyl, aryl, heterocyclyl or heteroaryl radical
20 optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or
25 (3) aryl, heteroaryl, heterocyclyl or C₃-C₆ cycloalkyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;
30 more preferably, each R₃₂ is independently
(1) hydrogen radicals;

(2) C₁-C₄ alkyl radical optionally substituted by phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl or trifluoromethyl radicals; or

(3) phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl or trifluoromethyl radicals;

more preferably, each R₃₂ is independently

(1) hydrogen radicals;

(2) C₁-C₄ alkyl radical or C₁-C₂ alkyl radical substituted by phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals; or

(3) phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals;

most preferably, R₃₂ is independently

(1) hydrogen or C₁-C₄ alkyl radical; or

(2) phenyl or heteroaryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals; and

wherein each R₃₃ is independently

(1) hydrogen radical; or

(2) alkyl radical optionally substituted by a radical of heterocycl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino,

alkanoylamino, alkoxycarbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano, alkyl or haloalkyl;

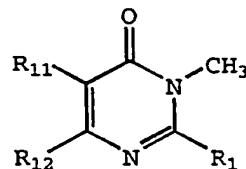
preferably, each R₃₃ is independently

- 5 (1) hydrogen radical; or
- (2) C₁-C₄ alkyl radical optionally substituted by a radical of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

more preferably, each R₃₃ is independently hydrogen or
15 C₁-C₄ alkyl radical; and most preferably, each R₃₃ is independently hydrogen or methyl radical.

The compounds of this invention may have in general several asymmetric centers and are typically depicted in
20 the form of racemic mixtures. This invention is intended to encompass racemic mixtures, partially racemic mixtures and separate enantiomers and diasteromers.

Compounds of interest include the following:



25

wherein R¹¹, R¹², and R¹ are one of the combinations given in the following table:

R ¹¹	R ¹²	R ¹
Phenyl	4-pyridyl	1-piperazinyl
4-fluorophenyl	4-pyridyl	1-piperazinyl
3-fluorophenyl	4-pyridyl	1-piperazinyl
2-fluorophenyl	4-pyridyl	1-piperazinyl
4-chlorophenyl	4-pyridyl	1-piperazinyl
3-chlorophenyl	4-pyridyl	1-piperazinyl

2-chlorophenyl	4-pyridyl	1-piperazinyl
4-tolyl	4-pyridyl	1-piperazinyl
3-tolyl	4-pyridyl	1-piperazinyl
2-tolyl	4-pyridyl	1-piperazinyl
4-trifluoro-methylphenyl	4-pyridyl	1-piperazinyl
3-trifluoro-methylphenyl	4-pyridyl	1-piperazinyl
2,6-dichlorophenyl	4-pyridyl	1-piperazinyl
2,6-dimethylphenyl	4-pyridyl	1-piperazinyl
3,4-dichlorophenyl	4-pyridyl	1-piperazinyl
3,4-dimethylphenyl	4-pyridyl	1-piperazinyl
2,4-dichlorophenyl	4-pyridyl	1-piperazinyl
2,4-dimethylphenyl	4-pyridyl	1-piperazinyl
Phenyl	2-amino-4-pyridyl	1-piperazinyl
4-fluorophenyl	2-amino-4-pyridyl	1-piperazinyl
3-fluorophenyl	2-amino-4-pyridyl	1-piperazinyl
2-fluorophenyl	2-amino-4-pyridyl	1-piperazinyl
4-chlorophenyl	2-amino-4-pyridyl	1-piperazinyl
3-chlorophenyl	2-amino-4-pyridyl	1-piperazinyl
2-chlorophenyl	2-amino-4-pyridyl	1-piperazinyl
4-tolyl	2-amino-4-pyridyl	1-piperazinyl
3-tolyl	2-amino-4-pyridyl	1-piperazinyl
2-tolyl	2-amino-4-pyridyl	1-piperazinyl
4-trifluoro-methylphenyl	2-amino-4-pyridyl	1-piperazinyl
3-trifluoro-methylphenyl	2-amino-4-pyridyl	1-piperazinyl
2,6-dichlorophenyl	2-amino-4-pyridyl	1-piperazinyl
2,6-dimethylphenyl	2-amino-4-pyridyl	1-piperazinyl
3,4-dichlorophenyl	2-amino-4-pyridyl	1-piperazinyl
3,4-dimethylphenyl	2-amino-4-pyridyl	1-piperazinyl
2,4-dichlorophenyl	2-amino-4-pyridyl	1-piperazinyl

2,4-dimethylphenyl	2-amino-4-pyridyl	1-piperazinyl
Phenyl	2-acetamido-4-pyridyl	1-piperazinyl
4-fluorophenyl	2-acetamido-4-pyridyl	1-piperazinyl
3-fluorophenyl	2-acetamido-4-pyridyl	1-piperazinyl
2-fluorophenyl	2-acetamido-4-pyridyl	1-piperazinyl
4-chlorophenyl	2-acetamido-4-pyridyl	1-piperazinyl
3-chlorophenyl	2-acetamido-4-pyridyl	1-piperazinyl
2-chlorophenyl	2-acetamido-4-pyridyl	1-piperazinyl
4-tolyl	2-acetamido-4-pyridyl	1-piperazinyl
3-tolyl	2-acetamido-4-pyridyl	1-piperazinyl
2-tolyl	2-acetamido-4-pyridyl	1-piperazinyl
4-trifluoro-methylphenyl	2-acetamido-4-pyridyl	1-piperazinyl
3-trifluoro-methylphenyl	2-acetamido-4-pyridyl	1-piperazinyl
2,6-dichlorophenyl	2-acetamido-4-pyridyl	1-piperazinyl
2,6-dimethylphenyl	2-acetamido-4-pyridyl	1-piperazinyl
3,4-dichlorophenyl	2-acetamido-4-pyridyl	1-piperazinyl
3,4-dimethylphenyl	2-acetamido-4-pyridyl	1-piperazinyl
2,4-dichlorophenyl	2-acetamido-4-pyridyl	1-piperazinyl
2,4-dimethylphenyl	2-acetamido-4-pyridyl	1-piperazinyl
Phenyl	2-amino-4-pyrimidinyl	1-piperazinyl
4-fluorophenyl	2-amino-4-pyrimidinyl	1-piperazinyl
3-fluorophenyl	2-amino-4-pyrimidinyl	1-piperazinyl
2-fluorophenyl	2-amino-4-pyrimidinyl	1-piperazinyl
4-chlorophenyl	2-amino-4-pyrimidinyl	1-piperazinyl
3-chlorophenyl	2-amino-4-pyrimidinyl	1-piperazinyl
2-chlorophenyl	2-amino-4-pyrimidinyl	1-piperazinyl
4-tolyl	2-amino-4-pyrimidinyl	1-piperazinyl

3-tolyl	2-amino-4-pyrimidinyl	1-piperazinyl
2-tolyl	2-amino-4-pyrimidinyl	1-piperazinyl
4-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	1-piperazinyl
3-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	1-piperazinyl
2,6-dichlorophenyl	2-amino-4-pyrimidinyl	1-piperazinyl
2,6-dimethylphenyl	2-amino-4-pyrimidinyl	1-piperazinyl
3,4-dichlorophenyl	2-amino-4-pyrimidinyl	1-piperazinyl
3,4-dimethylphenyl	2-amino-4-pyrimidinyl	1-piperazinyl
2,4-dichlorophenyl	2-amino-4-pyrimidinyl	1-piperazinyl
2,4-dimethylphenyl	2-amino-4-pyrimidinyl	1-piperazinyl
Phenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
4-fluorophenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
3-fluorophenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
2-fluorophenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
4-chlorophenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
3-chlorophenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
2-chlorophenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
4-tolyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
3-tolyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
2-tolyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
4-trifluoro-methylphenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
3-trifluoro-methylphenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
2,6-dichlorophenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
2,6-dimethylphenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
3,4-dichlorophenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
3,4-dimethylphenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino
2,4-dichlorophenyl	4-pyridyl	2-(2-chlorophenyl)ethylamino

2, 4-dimethyl phenyl	4-pyridyl	2-(2-chlorophenyl) ethylamino
4-fluorophenyl	4-pyridyl	3-(3-fluorophenyl) propylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	3-(3-fluorophenyl) propylamino
benzyl	4-pyridyl	3-phenylpropylamino
benzyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
2-thienyl	4-pyridyl	3-phenylpropylamino
2-thienyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
cyclohexyl	4-pyridyl	3-phenylpropylamino
cyclohexyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
tert-butyl	4-pyridyl	3-phenylpropylamino
tert-butyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
4-fluorophenyl	4-piperidinyl	3-phenylpropylamino
4-fluorophenyl	4-piperidinyl	2-(4-fluorophenyl) ethylamino
4-fluorophenyl	4-pyranyl	3-phenylpropylamino
4-fluorophenyl	4-pyranyl	2-(4-fluorophenyl) ethylamino
Phenyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
4-fluorophenyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
3-fluorophenyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
2-fluorophenyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
4-chlorophenyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
3-chlorophenyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
2-chlorophenyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
4-tolyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
3-tolyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
2-tolyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
4-trifluoro-methylphenyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
3-trifluoro-methylphenyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
2, 6-dichlorophenyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino
2, 6-dimethyl phenyl	2-amino-4-pyridyl	2-(2-chlorophenyl) ethylamino

3, 4-dichlorophenyl	2-amino-4-pyridyl	2-(2-chlorophenyl)ethylamino
3, 4-dimethylphenyl	2-amino-4-pyridyl	2-(2-chlorophenyl)ethylamino
2, 4-dichlorophenyl	2-amino-4-pyridyl	2-(2-chlorophenyl)ethylamino
2, 4-dimethylphenyl	2-amino-4-pyridyl	2-(2-chlorophenyl)ethylamino
Phenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
4-fluorophenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
3-fluorophenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
2-fluorophenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
4-chlorophenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
3-chlorophenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
2-chlorophenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
4-tolyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
3-tolyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
2-tolyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
4-trifluoro-methylphenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
3-trifluoro-methylphenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
2, 6-dichlorophenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
2, 6-dimethylphenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
3, 4-dichlorophenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
3, 4-dimethylphenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
2, 4-dichlorophenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
2, 4-dimethylphenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl)ethylamino
Phenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
3-fluorophenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
2-fluorophenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
4-chlorophenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino

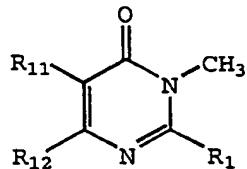
3-chlorophenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
2-chlorophenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
4-tolyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
3-tolyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
2-tolyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
4-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
3-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
2,6-dichlorophenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
2,6-dimethylphenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
3,4-dichlorophenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
3,4-dimethylphenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
2,4-dichlorophenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
2,4-dimethylphenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl)ethylamino
Phenyl	4-pyridyl	3-imidazolylpropylamino
4-fluorophenyl	4-pyridyl	3-imidazolylpropylamino
3-fluorophenyl	4-pyridyl	3-imidazolylpropylamino
2-fluorophenyl	4-pyridyl	3-imidazolylpropylamino
4-chlorophenyl	4-pyridyl	3-imidazolylpropylamino
3-chlorophenyl	4-pyridyl	3-imidazolylpropylamino
2-chlorophenyl	4-pyridyl	3-imidazolylpropylamino
4-tolyl	4-pyridyl	3-imidazolylpropylamino
3-tolyl	4-pyridyl	3-imidazolylpropylamino
2-tolyl	4-pyridyl	3-imidazolylpropylamino
4-trifluoro-methylphenyl	4-pyridyl	3-imidazolylpropylamino
3-trifluoro-methylphenyl	4-pyridyl	3-imidazolylpropylamino
2,6-dichlorophenyl	4-pyridyl	3-imidazolylpropylamino
2,6-dimethylphenyl	4-pyridyl	3-imidazolylpropylamino
3,4-dichlorophenyl	4-pyridyl	3-imidazolylpropylamino
3,4-dimethylphenyl	4-pyridyl	3-imidazolylpropylamino
2,4-dichlorophenyl	4-pyridyl	3-imidazolylpropylamino
2,4-dimethylphenyl	4-pyridyl	3-imidazolylpropylamino
Phenyl	2-amino-4-pyridyl	3-imidazolylpropylamino

4-fluorophenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
3-fluorophenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
2-fluorophenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
4-chlorophenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
3-chlorophenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
2-chlorophenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
4-tolyl	2-amino-4-pyridyl	3-imidazolylpropylamino
3-tolyl	2-amino-4-pyridyl	3-imidazolylpropylamino
2-tolyl	2-amino-4-pyridyl	3-imidazolylpropylamino
4-trifluoro-methylphenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
3-trifluoro-methylphenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
2,6-dichlorophenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
2,6-dimethylphenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
3,4-dichlorophenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
3,4-dimethylphenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
2,4-dichlorophenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
2,4-dimethylphenyl	2-amino-4-pyridyl	3-imidazolylpropylamino
Phenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
4-fluorophenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
3-fluorophenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
2-fluorophenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
4-chlorophenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
3-chlorophenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
2-chlorophenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
4-tolyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
3-tolyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
2-tolyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino

4-trifluoro-methylphenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
3-trifluoro-methylphenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
2,6-dichlorophenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
2,6-dimethylphenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
3,4-dichlorophenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
3,4-dimethylphenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
2,4-dichlorophenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
2,4-dimethylphenyl	2-acetamido-4-pyridyl	3-imidazolylpropylamino
Phenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
3-fluorophenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
2-fluorophenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
4-chlorophenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
3-chlorophenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
2-chlorophenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
4-tolyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
3-tolyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
2-tolyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
4-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
3-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
2,6-dichlorophenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
2,6-dimethylphenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
3,4-dichlorophenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
3,4-dimethylphenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
2,4-dichlorophenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
2,4-dimethylphenyl	2-amino-4-pyrimidinyl	3-imidazolylpropylamino
4-fluorophenyl	4-pyridyl	2-(2-chlorophenyl-1-methyl)ethylamino

4-fluorophenyl	2-acetamido-4-pyridyl	2-(2-chlorophenyl-1-methyl)ethyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	2-(2-chlorophenyl-1-methyl)ethyl)amino
3-fluorophenyl	4-pyridyl	(S)-tetrahydroisoquinol-3-ylmethylenamino
2-fluorophenyl	2-amino-4-pyridyl	(S)-3-benzylpiperazinyl
3-chlorophenyl	2-acetamido-4-pyridyl	(S)-2-N-isopropylamino-3-phenylpropylamino
2-chlorophenyl	2-amino-4-pyrimidinyl	(S)-2-N-glycylamino-3-phenylpropylamino
4-tolyl	4-pyridyl	(S)-2-amino-3-phenylpropylamino
3-tolyl	2-amino-4-pyridyl	(R)-2-amino-3-phenylpropylamino
2-tolyl	2-acetamido-4-pyridyl	3-amino-3-phenylpropylamino
4-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	(S)-2-amino-3-(2-fluorophenyl)propylamino
3-trifluoro-methylphenyl	4-pyridyl	(S)-2-amino-3-(2-methylphenyl)propylamino
2,6-dichlorophenyl	2-amino-4-pyridyl	3-amino-3-(2-fluorophenyl)propylamino
2,6-dimethylphenyl	2-acetamido-4-pyridyl	3-amino-3-(2-methylphenyl)propylamino
3,4-dichlorophenyl	2-amino-4-pyrimidinyl	2-amino-2-methyl-3-phenylpropylamino
3,4-dimethylphenyl	4-pyridyl	3-amino-2-methyl-3-phenylpropylamino
3-fluorophenyl	2-amino-4-pyridyl	(S)-2-amino-3-phenylpropylamino
2-fluorophenyl	2-acetamido-4-pyridyl	(S)-2-amino-3-(2-fluorophenyl)propylamino
3-chlorophenyl	2-amino-4-pyrimidinyl	(S)-2-amino-3-(2-methylphenyl)propylamino
2-chlorophenyl	4-pyridyl	(S)-2-N-isopropylamino-3-phenylpropylamino
4-tolyl	2-amino-4-pyridyl	(S)-2-N-glycylamino-3-phenylpropylamino
3-tolyl	2-acetamido-4-pyridyl	2-amino-2-methyl-3-phenylpropylamino
2-tolyl	2-amino-4-pyrimidinyl	(R)-2-amino-3-phenylpropylamino
4-trifluoro-methylphenyl	4-pyridyl	3-amino-3-phenylpropylamino
3-trifluoro-methylphenyl	2-amino-4-pyridyl	3-amino-3-(2-fluorophenyl)propylamino
2,6-dichlorophenyl	2-acetamido-4-pyridyl	3-amino-3-(2-methylphenyl)propylamino
2,6-dimethylphenyl	2-amino-4-pyrimidinyl	3-amino-2-methyl-3-phenylpropylamino
3,4-dichlorophenyl	4-pyridyl	(S)-tetrahydroisoquinol-3-ylmethylenamino

3, 4-dimethyl phenyl and	4-pyridyl	(S)-3-benzylpiperazinyl
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wherein R^{11} , R^{12} , and R^1 are one of the combinations given in the following table:

R^{11}	R^{12}	R^1
Phenyl	4-pyridyl	4-pyridyl
4-fluorophenyl	4-pyridyl	4-pyridyl
3-fluorophenyl	4-pyridyl	4-pyridyl
2-fluorophenyl	4-pyridyl	4-pyridyl
4-chlorophenyl	4-pyridyl	4-pyridyl
3-chlorophenyl	4-pyridyl	4-pyridyl
2-chlorophenyl	4-pyridyl	4-pyridyl
4-tolyl	4-pyridyl	4-pyridyl
3-tolyl	4-pyridyl	4-pyridyl
2-tolyl	4-pyridyl	4-pyridyl
4-trifluoro-methylphenyl	4-pyridyl	4-pyridyl
3-trifluoro-methylphenyl	4-pyridyl	4-pyridyl
2, 6-dichlorophenyl	4-pyridyl	4-pyridyl
2, 6-dimethyl phenyl	4-pyridyl	4-pyridyl
3, 4-dichlorophenyl	4-pyridyl	4-pyridyl
3, 4-dimethyl phenyl	4-pyridyl	4-pyridyl
2, 4-dichlorophenyl	4-pyridyl	4-pyridyl
2, 4-dimethyl phenyl	4-pyridyl	4-pyridyl
Phenyl	2-amino-4-pyridyl	4-pyridyl
4-fluorophenyl	2-amino-4-pyridyl	4-pyridyl
3-fluorophenyl	2-amino-4-pyridyl	4-pyridyl
2-fluorophenyl	2-amino-4-pyridyl	4-pyridyl
4-chlorophenyl	2-amino-4-pyridyl	4-pyridyl
3-chlorophenyl	2-amino-4-pyridyl	4-pyridyl

2-chlorophenyl	2-amino-4-pyridyl	4-pyridyl
4-tolyl	2-amino-4-pyridyl	4-pyridyl
3-tolyl	2-amino-4-pyridyl	4-pyridyl
2-tolyl	2-amino-4-pyridyl	4-pyridyl
4-trifluoro-methylphenyl	2-amino-4-pyridyl	4-pyridyl
3-trifluoro-methylphenyl	2-amino-4-pyridyl	4-pyridyl
2,6-dichlorophenyl	2-amino-4-pyridyl	4-pyridyl
2,6-dimethylphenyl	2-amino-4-pyridyl	4-pyridyl
3,4-dichlorophenyl	2-amino-4-pyridyl	4-pyridyl
3,4-dimethylphenyl	2-amino-4-pyridyl	4-pyridyl
2,4-dichlorophenyl	2-amino-4-pyridyl	4-pyridyl
2,4-dimethylphenyl	2-amino-4-pyridyl	4-pyridyl
Phenyl	2-acetamido-4-pyridyl	4-pyridyl
4-fluorophenyl	2-acetamido-4-pyridyl	4-pyridyl
3-fluorophenyl	2-acetamido-4-pyridyl	4-pyridyl
2-fluorophenyl	2-acetamido-4-pyridyl	4-pyridyl
4-chlorophenyl	2-acetamido-4-pyridyl	4-pyridyl
3-chlorophenyl	2-acetamido-4-pyridyl	4-pyridyl
2-chlorophenyl	2-acetamido-4-pyridyl	4-pyridyl
4-tolyl	2-acetamido-4-pyridyl	4-pyridyl
3-tolyl	2-acetamido-4-pyridyl	4-pyridyl
2-tolyl	2-acetamido-4-pyridyl	4-pyridyl
4-trifluoro-methylphenyl	2-acetamido-4-pyridyl	4-pyridyl
3-trifluoro-methylphenyl	2-acetamido-4-pyridyl	4-pyridyl
2,6-dichlorophenyl	2-acetamido-4-pyridyl	4-pyridyl
2,6-dimethylphenyl	2-acetamido-4-pyridyl	4-pyridyl
3,4-dichlorophenyl	2-acetamido-4-pyridyl	4-pyridyl

3, 4-dimethyl phenyl	2-acetamido-4-pyridyl	4-pyridyl
2, 4-dichlorophenyl	2-acetamido-4-pyridyl	4-pyridyl
2, 4-dimethyl phenyl	2-acetamido-4-pyridyl	4-pyridyl
Phenyl	2-amino-4-pyrimidinyl	4-pyridyl
4-fluorophenyl	2-amino-4-pyrimidinyl	4-pyridyl
3-fluorophenyl	2-amino-4-pyrimidinyl	4-pyridyl
2-fluorophenyl	2-amino-4-pyrimidinyl	4-pyridyl
4-chlorophenyl	2-amino-4-pyrimidinyl	4-pyridyl
3-chlorophenyl	2-amino-4-pyrimidinyl	4-pyridyl
2-chlorophenyl	2-amino-4-pyrimidinyl	4-pyridyl
4-tolyl	2-amino-4-pyrimidinyl	4-pyridyl
3-tolyl	2-amino-4-pyrimidinyl	4-pyridyl
2-tolyl	2-amino-4-pyrimidinyl	4-pyridyl
4-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	4-pyridyl
3-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	4-pyridyl
2, 6-dichlorophenyl	2-amino-4-pyrimidinyl	4-pyridyl
2, 6-dimethyl phenyl	2-amino-4-pyrimidinyl	4-pyridyl
3, 4-dichlorophenyl	2-amino-4-pyrimidinyl	4-pyridyl
3, 4-dimethyl phenyl	2-amino-4-pyrimidinyl	4-pyridyl
2, 4-dichlorophenyl	2-amino-4-pyrimidinyl	4-pyridyl
2, 4-dimethyl phenyl	2-amino-4-pyrimidinyl	4-pyridyl
Phenyl	4-pyridyl	4-methyl sulfinylphenyl
4-fluorophenyl	4-pyridyl	4-methyl sulfinylphenyl
3-fluorophenyl	4-pyridyl	4-methyl sulfinylphenyl
2-fluorophenyl	4-pyridyl	4-methyl sulfinylphenyl
4-chlorophenyl	4-pyridyl	4-methyl sulfinylphenyl
3-chlorophenyl	4-pyridyl	4-methyl sulfinylphenyl
2-chlorophenyl	4-pyridyl	4-methyl sulfinylphenyl
4-tolyl	4-pyridyl	4-methyl sulfinylphenyl
3-tolyl	4-pyridyl	4-methyl sulfinylphenyl
2-tolyl	4-pyridyl	4-methyl sulfinylphenyl
4-trifluoro-methylphenyl	4-pyridyl	4-methyl sulfinylphenyl

<u>3-trifluoro-</u> <u>methylphenyl</u>	4-pyridyl	4-methyl sulfinylphenyl
<u>2,6-</u> <u>dichlorophenyl</u>	4-pyridyl	4-methyl sulfinylphenyl
<u>2,6-dimethyl</u> <u>phenyl</u>	4-pyridyl	4-methyl sulfinylphenyl
<u>3,4-</u> <u>dichlorophenyl</u>	4-pyridyl	4-methyl sulfinylphenyl
<u>3,4-dimethyl</u> <u>phenyl</u>	4-pyridyl	4-methyl sulfinylphenyl
<u>2,4-</u> <u>dichlorophenyl</u>	4-pyridyl	4-methyl sulfinylphenyl
<u>2,4-dimethyl</u> <u>phenyl</u>	4-pyridyl	4-methyl sulfinylphenyl
<u>Phenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>4-fluorophenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>3-fluorophenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>2-fluorophenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>4-chlorophenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>3-chlorophenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>2-chlorophenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>4-tolyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>3-tolyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>2-tolyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>4-trifluoro-</u> <u>methylphenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>3-trifluoro-</u> <u>methylphenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>2,6-</u> <u>dichlorophenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>2,6-dimethyl</u> <u>phenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>3,4-</u> <u>dichlorophenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>3,4-dimethyl</u> <u>phenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>2,4-</u> <u>dichlorophenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>2,4-dimethyl</u> <u>phenyl</u>	<u>2-amino-4-</u> <u>pyridyl</u>	4-methyl sulfinylphenyl
<u>Phenyl</u>	<u>2-acetamido-</u> <u>4-pyridyl</u>	4-methyl sulfinylphenyl
<u>4-fluorophenyl</u>	<u>2-acetamido-</u> <u>4-pyridyl</u>	4-methyl sulfinylphenyl

3-fluorophenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
2-fluorophenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
4-chlorophenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
3-chlorophenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
2-chlorophenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
4-tolyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
3-tolyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
2-tolyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
4-trifluoro-methylphenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
3-trifluoro-methylphenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
2,6-dichlorophenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
2,6-dimethylphenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
3,4-dichlorophenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
3,4-dimethylphenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
2,4-dichlorophenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
2,4-dimethylphenyl	2-acetamido-4-pyridyl	4-methyl sulfinylphenyl
Phenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
4-fluorophenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
3-fluorophenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
2-fluorophenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
4-chlorophenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
3-chlorophenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
2-chlorophenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
4-tolyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
3-tolyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
2-tolyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
4-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl

3-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
2,6-dichlorophenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
2,6-dimethyl phenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
3,4-dichlorophenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
3,4-dimethyl phenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
2,4-dichlorophenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
2,4-dimethyl phenyl	2-amino-4-pyrimidinyl	4-methyl sulfinylphenyl
Phenyl	4-pyridyl	2,6-dichlorobenzyl
4-fluorophenyl	4-pyridyl	2,6-dichlorobenzyl
3-fluorophenyl	4-pyridyl	2,6-dichlorobenzyl
2-fluorophenyl	4-pyridyl	2,6-dichlorobenzyl
4-chlorophenyl	4-pyridyl	2,6-dichlorobenzyl
3-chlorophenyl	4-pyridyl	2,6-dichlorobenzyl
2-chlorophenyl	4-pyridyl	2,6-dichlorobenzyl
4-tolyl	4-pyridyl	2,6-dichlorobenzyl
3-tolyl	4-pyridyl	2,6-dichlorobenzyl
2-tolyl	4-pyridyl	2,6-dichlorobenzyl
4-trifluoro-methylphenyl	4-pyridyl	2,6-dichlorobenzyl
3-trifluoro-methylphenyl	4-pyridyl	2,6-dichlorobenzyl
2,6-dichlorophenyl	4-pyridyl	2,6-dichlorobenzyl
2,6-dimethyl phenyl	4-pyridyl	2,6-dichlorobenzyl
3,4-dichlorophenyl	4-pyridyl	2,6-dichlorobenzyl
3,4-dimethyl phenyl	4-pyridyl	2,6-dichlorobenzyl
2,4-dichlorophenyl	4-pyridyl	2,6-dichlorobenzyl
2,4-dimethyl phenyl	4-pyridyl	2,6-dichlorobenzyl
Phenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
4-fluorophenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
3-fluorophenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
2-fluorophenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
4-chlorophenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
3-chlorophenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
2-chlorophenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl

4-tolyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
3-tolyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
2-tolyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
4-trifluoro-methylphenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
3-trifluoro-methylphenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
2,6-dichlorophenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
2,6-dimethylphenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
3,4-dichlorophenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
3,4-dimethylphenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
2,4-dichlorophenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
2,4-dimethylphenyl	2-amino-4-pyridyl	2,6-dichlorobenzyl
Phenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
4-fluorophenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
3-fluorophenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
2-fluorophenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
4-chlorophenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
3-chlorophenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
2-chlorophenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
4-tolyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
3-tolyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
2-tolyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
4-trifluoro-methylphenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
3-trifluoro-methylphenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
2,6-dichlorophenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
2,6-dimethylphenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
3,4-dichlorophenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
3,4-dimethylphenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl

2,4-dichlorophenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
2,4-dimethylphenyl	2-acetamido-4-pyridyl	2,6-dichlorobenzyl
Phenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
4-fluorophenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
3-fluorophenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
2-fluorophenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
4-chlorophenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
3-chlorophenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
2-chlorophenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
4-tolyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
3-tolyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
2-tolyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
4-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
3-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
2,6-dichlorophenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
2,6-dimethylphenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
3,4-dichlorophenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
3,4-dimethylphenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
2,4-dichlorophenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
2,4-dimethylphenyl	2-amino-4-pyrimidinyl	2,6-dichlorobenzyl
Phenyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
4-fluorophenyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
3-fluorophenyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
2-fluorophenyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
4-chlorophenyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
3-chlorophenyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
2-chlorophenyl	4-pyridyl	2-(4-fluorophenyl)ethylamino

4-tolyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
3-tolyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
2-tolyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
4-trifluoro-methylphenyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
3-trifluoro-methylphenyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
2,6-dichlorophenyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
2,6-dimethylphenyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
3,4-dichlorophenyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
3,4-dimethylphenyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
2,4-dichlorophenyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
2,4-dimethylphenyl	4-pyridyl	2-(4-fluorophenyl) ethylamino
Phenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
4-fluorophenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
3-fluorophenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
2-fluorophenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
4-chlorophenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
3-chlorophenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
2-chlorophenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
4-tolyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
3-tolyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
2-tolyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
4-trifluoro-methylphenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
3-trifluoro-methylphenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
2,6-dichlorophenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
2,6-dimethylphenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
3,4-dichlorophenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino
3,4-dimethylphenyl	2-amino-4-pyridyl	2-(4-fluorophenyl) ethylamino

2,4-dichlorophenyl	2-amino-4-pyridyl	2-(4-fluorophenyl)ethylamino
2,4-dimethylphenyl	2-amino-4-pyridyl	2-(4-fluorophenyl)ethylamino
Phenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
4-fluorophenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
3-fluorophenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
2-fluorophenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
4-chlorophenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
3-chlorophenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
2-chlorophenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
4-tolyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
3-tolyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
2-tolyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
4-trifluoro-methylphenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
3-trifluoro-methylphenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
2,6-dichlorophenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
2,6-dimethylphenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
3,4-dichlorophenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
3,4-dimethylphenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
2,4-dichlorophenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
2,4-dimethylphenyl	2-acetamido-4-pyridyl	2-(4-fluorophenyl)ethylamino
Phenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
3-fluorophenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
2-fluorophenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
4-chlorophenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
3-chlorophenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
2-chlorophenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino

4-tolyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
3-tolyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
2-tolyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
4-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
3-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
2,6-dichlorophenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
2,6-dimethylphenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
3,4-dichlorophenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
3,4-dimethylphenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
2,4-dichlorophenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
2,4-dimethylphenyl	2-amino-4-pyrimidinyl	2-(4-fluorophenyl)ethylamino
Phenyl	4-pyridyl	3-phenyl-propylamino
4-fluorophenyl	4-pyridyl	3-phenyl-propylamino
3-fluorophenyl	4-pyridyl	3-phenyl-propylamino
2-fluorophenyl	4-pyridyl	3-phenyl-propylamino
4-chlorophenyl	4-pyridyl	3-phenyl-propylamino
3-chlorophenyl	4-pyridyl	3-phenyl-propylamino
2-chlorophenyl	4-pyridyl	3-phenyl-propylamino
4-tolyl	4-pyridyl	3-phenyl-propylamino
3-tolyl	4-pyridyl	3-phenyl-propylamino
2-tolyl	4-pyridyl	3-phenyl-propylamino
4-trifluoro-methylphenyl	4-pyridyl	3-phenyl-propylamino
3-trifluoro-methylphenyl	4-pyridyl	3-phenyl-propylamino
2,6-dichlorophenyl	4-pyridyl	3-phenyl-propylamino
2,6-dimethylphenyl	4-pyridyl	3-phenyl-propylamino
3,4-dichlorophenyl	4-pyridyl	3-phenyl-propylamino
3,4-dimethylphenyl	4-pyridyl	3-phenyl-propylamino
2,4-dichlorophenyl	4-pyridyl	3-phenyl-propylamino
2,4-dimethylphenyl	4-pyridyl	3-phenyl-propylamino
Phenyl	2-amino-4-pyridyl	3-phenyl-propylamino
4-fluorophenyl	2-amino-4-pyridyl	3-phenyl-propylamino
3-fluorophenyl	2-amino-4-pyridyl	3-phenyl-propylamino

2-fluorophenyl	2-amino-4-pyridyl	3-phenyl-propylamino
4-chlorophenyl	2-amino-4-pyridyl	3-phenyl-propylamino
3-chlorophenyl	2-amino-4-pyridyl	3-phenyl-propylamino
2-chlorophenyl	2-amino-4-pyridyl	3-phenyl-propylamino
4-tolyl	2-amino-4-pyridyl	3-phenyl-propylamino
3-tolyl	2-amino-4-pyridyl	3-phenyl-propylamino
2-tolyl	2-amino-4-pyridyl	3-phenyl-propylamino
4-trifluoro-methylphenyl	2-amino-4-pyridyl	3-phenyl-propylamino
3-trifluoro-methylphenyl	2-amino-4-pyridyl	3-phenyl-propylamino
2,6-dichlorophenyl	2-amino-4-pyridyl	3-phenyl-propylamino
2,6-dimethylphenyl	2-amino-4-pyridyl	3-phenyl-propylamino
3,4-dichlorophenyl	2-amino-4-pyridyl	3-phenyl-propylamino
3,4-dimethylphenyl	2-amino-4-pyridyl	3-phenyl-propylamino
2,4-dichlorophenyl	2-amino-4-pyridyl	3-phenyl-propylamino
2,4-dimethylphenyl	2-amino-4-pyridyl	3-phenyl-propylamino
Phenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
4-fluorophenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
3-fluorophenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
2-fluorophenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
4-chlorophenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
3-chlorophenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
2-chlorophenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
4-tolyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
3-tolyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
2-tolyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
4-trifluoro-methylphenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
3-trifluoro-methylphenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino

2,6-dichlorophenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
2,6-dimethylphenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
3,4-dichlorophenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
3,4-dimethylphenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
2,4-dichlorophenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
2,4-dimethylphenyl	2-acetamido-4-pyridyl	3-phenyl-propylamino
Phenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
3-fluorophenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
2-fluorophenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
4-chlorophenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
3-chlorophenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
2-chlorophenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
4-tolyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
3-tolyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
2-tolyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
4-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
3-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
2,6-dichlorophenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
2,6-dimethylphenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
3,4-dichlorophenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
3,4-dimethylphenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
2,4-dichlorophenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
2,4-dimethylphenyl	2-amino-4-pyrimidinyl	3-phenyl-propylamino
Phenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
4-fluorophenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
3-fluorophenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino

2-fluorophenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
4-chlorophenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
3-chlorophenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
2-chlorophenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
4-tolyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
3-tolyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
2-tolyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
4-trifluoro-methylphenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
3-trifluoro-methylphenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
2,6-dichlorophenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
2,6-dimethylphenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
3,4-dichlorophenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
3,4-dimethylphenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
2,4-dichlorophenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
2,4-dimethylphenyl	4-pyridyl	(1-methyl-3-phenyl)propylamino
Phenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
4-fluorophenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
3-fluorophenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
2-fluorophenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
4-chlorophenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
3-chlorophenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
2-chlorophenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
4-tolyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
3-tolyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
2-tolyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
4-trifluoro-methylphenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
3-trifluoro-methylphenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino

2,6-dichlorophenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
2,6-dimethylphenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
3,4-dichlorophenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
3,4-dimethylphenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
2,4-dichlorophenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
2,4-dimethylphenyl	2-amino-4-pyridyl	(1-methyl-3-phenyl)propylamino
Phenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
4-fluorophenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
3-fluorophenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
2-fluorophenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
4-chlorophenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
3-chlorophenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
2-chlorophenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
4-tolyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
3-tolyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
2-tolyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
4-trifluoro-methylphenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
3-trifluoro-methylphenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
2,6-dichlorophenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
2,6-dimethylphenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
3,4-dichlorophenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
3,4-dimethylphenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
2,4-dichlorophenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
2,4-dimethylphenyl	2-acetamido-4-pyridyl	(1-methyl-3-phenyl)propylamino
Phenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
3-fluorophenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino

2-fluorophenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
4-chlorophenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
3-chlorophenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
2-chlorophenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
4-tolyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
3-tolyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
2-tolyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
4-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
3-trifluoro-methylphenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
2,6-dichlorophenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
2,6-dimethylphenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
3,4-dichlorophenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
3,4-dimethylphenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
2,4-dichlorophenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
2,4-dimethylphenyl	2-amino-4-pyrimidinyl	(1-methyl-3-phenyl)propylamino
4-fluorophenyl	4-pyridyl	4-fluorobenzylamino
4-fluorophenyl	2-acetamido-4-pyridyl	4-fluorobenzylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	4-fluorobenzylamino
4-fluorophenyl	4-pyridyl	(2-(4-fluorophenyl)-1-methyl-ethyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(2-(4-fluorophenyl)-1-methyl-ethyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(2-(4-fluorophenyl)-1-methyl-ethyl)amino
4-fluorophenyl	4-pyridyl	(1,1-dimethyl-2-(4-fluorophenyl)-ethyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(1,1-dimethyl-2-(4-fluorophenyl)-ethyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(1,1-dimethyl-2-(4-fluorophenyl)-ethyl)amino
4-fluorophenyl	4-pyridyl	2-(4-fluorophenyl)-2-methyl-ethylamino
4-fluorophenyl	2-acetamido-4-pyridyl	(2-(4-fluorophenyl)-2-methyl-ethyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(2-(4-fluorophenyl)-2-methyl-ethyl)amino

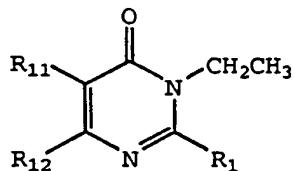
4-fluorophenyl	4-pyridyl	(2-methyl-2-phenylethyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(2-methyl-2-phenylethyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(2-methyl-2-phenylethyl)amino
4-fluorophenyl	4-pyridyl	methyl-(2-phenylethyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	methyl-(2-phenylethyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	methyl-(2-phenylethyl)amino
4-fluorophenyl	4-pyridyl	(2-(4-trifluoromethylphenyl)ethyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(2-(4-trifluoromethylphenyl)ethyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(2-(4-trifluoromethylphenyl)ethyl)amino
4-fluorophenyl	4-pyridyl	2-(4-tolyl)ethylamino
4-fluorophenyl	2-acetamido-4-pyridyl	2-(4-tolyl)ethylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	2-(4-tolyl)ethylamino
4-fluorophenyl	4-pyridyl	(2-(3-fluorophenyl)ethyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(2-(3-fluorophenyl)ethyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(2-(3-fluorophenyl)ethyl)amino
4-fluorophenyl	4-pyridyl	(2-(2-fluorophenyl)ethyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(2-(2-fluorophenyl)ethyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(2-(2-fluorophenyl)ethyl)amino
4-fluorophenyl	4-pyridyl	methyl-(2-(2-pyridyl)ethyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	methyl-(2-(2-pyridyl)ethyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	methyl-(2-(2-pyridyl)ethyl)amino
4-fluorophenyl	4-pyridyl	(1,1-dimethyl-3-phenylpropyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(1,1-dimethyl-3-phenylpropyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(1,1-dimethyl-3-phenylpropyl)amino
4-fluorophenyl	4-pyridyl	(3-(4-fluorophenyl)-propyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(3-(4-fluorophenyl)-propyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(3-(4-fluorophenyl)-propyl)amino

4-fluorophenyl	4-pyridyl	(3-(4-fluorophenyl)-1-methyl-propyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(3-(4-fluorophenyl)-1-methyl-propyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(3-(4-fluorophenyl)-1-methyl-propyl)amino
4-fluorophenyl	4-pyridyl	(1,1-dimethyl-3-(4-fluoro phenyl)-propyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(1,1-dimethyl-3-(4-fluoro phenyl)-propyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(1,1-dimethyl-3-(4-fluoro phenyl)-propyl)amino
4-fluorophenyl	4-pyridyl	(3-(2-fluorophenyl)-propyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(3-(2-fluorophenyl)-propyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(3-(2-fluorophenyl)-propyl)amino
4-fluorophenyl	4-pyridyl	(3-methyl-3-phenyl-propyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(3-methyl-3-phenyl-propyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(3-methyl-3-phenyl-propyl)amino
4-fluorophenyl	4-pyridyl	(2-methyl-3-phenyl-propyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(2-methyl-3-phenyl-propyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(2-methyl-3-phenyl-propyl)amino
4-fluorophenyl	4-pyridyl	(3,3-dimethylbutyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(3,3-dimethylbutyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(3,3-dimethylbutyl)amino
4-fluorophenyl	4-pyridyl	isoamylamino
4-fluorophenyl	2-acetamido-4-pyridyl	isoamylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	isoamylamino
4-fluorophenyl	4-pyridyl	amylamino
4-fluorophenyl	2-acetamido-4-pyridyl	amylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	amylamino
4-fluorophenyl	4-pyridyl	(2,5-dimethyl)pentylamino
4-fluorophenyl	2-acetamido-4-pyridyl	(2,5-dimethyl)pentylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	(2,5-dimethyl)pentylamino
4-fluorophenyl	4-pyridyl	piperazinyl
4-fluorophenyl	2-acetamido-4-pyridyl	piperazinyl

4-fluorophenyl	2-amino-4-pyrimidinyl	piperazinyl
4-fluorophenyl	4-pyridyl	(3-(3-fluorophenyl)-propyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(3-(3-fluorophenyl)-propyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(3-(3-fluorophenyl)-propyl)amino
benzyl	4-pyridyl	3-phenylpropylamino
benzyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
2-thienyl	4-pyridyl	3-phenylpropylamino
2-thienyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
cyclohexyl	4-pyridyl	3-phenylpropylamino
cyclohexyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
tert-butyl	4-pyridyl	3-phenylpropylamino
tert-butyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
4-fluorophenyl	4-piperidinyl	3-phenylpropylamino
4-fluorophenyl	4-piperidinyl	2-(4-fluorophenyl)ethylamino
4-fluorophenyl	4-pyranyl	3-phenylpropylamino
4-fluorophenyl	4-pyranyl	2-(4-fluorophenyl)ethylamino
Phenyl	4-pyridyl	3-phenyl-2-amino-propylamino
4-fluorophenyl	4-pyridyl	3-phenyl-2-amino-propylamino
3-fluorophenyl	4-pyridyl	3-phenyl-2-amino-propylamino
2-fluorophenyl	4-pyridyl	3-phenyl-2-amino-propylamino
4-chlorophenyl	4-pyridyl	3-phenyl-2-amino-propylamino
3-chlorophenyl	4-pyridyl	3-phenyl-2-amino-propylamino
2-chlorophenyl	4-pyridyl	3-phenyl-2-amino-propylamino
4-tolyl	4-pyridyl	3-phenyl-2-amino-propylamino
3-tolyl	4-pyridyl	3-phenyl-2-amino-propylamino
2-tolyl	4-pyridyl	3-phenyl-2-amino-propylamino
4-trifluoro-methylphenyl	4-pyridyl	3-phenyl-2-amino-propylamino
3-trifluoro-methylphenyl	4-pyridyl	3-phenyl-2-amino-propylamino
2,6-dichlorophenyl	4-pyridyl	3-phenyl-2-amino-propylamino

<u>2,6-dimethyl phenyl</u>	4-pyridyl	3-phenyl-2-amino-propylamino
<u>3,4-dichlorophenyl</u>	4-pyridyl	3-phenyl-2-amino-propylamino
<u>3,4-dimethyl phenyl</u>	4-pyridyl	3-phenyl-2-amino-propylamino
<u>2,4-dichlorophenyl</u>	4-pyridyl	3-phenyl-2-amino-propylamino
<u>2,4-dimethyl phenyl</u>	4-pyridyl	3-phenyl-2-amino-propylamino
<u>Phenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>4-fluorophenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>3-fluorophenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>2-fluorophenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>4-chlorophenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>3-chlorophenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>2-chlorophenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>4-tolyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>3-tolyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>2-tolyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>4-trifluoro-methylphenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>3-trifluoro-methylphenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>2,6-dichlorophenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>2,6-dimethyl phenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>3,4-dichlorophenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>3,4-dimethyl phenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>2,4-dichlorophenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino
<u>2,4-dimethyl phenyl</u>	4-pyridyl	3-phenyl-3-amino-propylamino

and

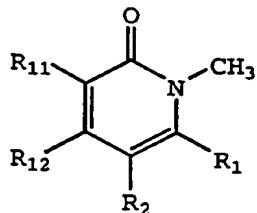


wherein R^{11} , R^{12} , and R^1 are one of the combinations given in the following table:

R^{11}	R^{12}	R^1
4-fluorophenyl	4-pyridyl	(2-(4-fluorophenyl)ethyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(2-(4-fluorophenyl)ethyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(2-(4-fluorophenyl)ethyl)amino
4-fluorophenyl	4-pyridyl	(3-phenylpropyl)amino
4-fluorophenyl	2-acetamido-4-pyridyl	(3-phenylpropyl)amino
4-fluorophenyl	2-amino-4-pyrimidinyl	(3-phenylpropyl)amino
4-fluorophenyl	4-pyridyl	(S)-2-amino-3-phenylpropylamino
4-fluorophenyl	2-acetamido-4-pyridyl	(S)-2-amino-3-phenylpropylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	(S)-2-amino-3-phenylpropylamino
4-fluorophenyl	4-pyridyl	3-amino-3-phenylpropylamino
4-fluorophenyl	2-acetamido-4-pyridyl	3-amino-3-phenylpropylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	3-amino-3-phenylpropylamino
4-fluorophenyl	4-pyridyl	3-amino-2-methyl-3-phenylpropylamino
4-fluorophenyl	2-acetamido-4-pyridyl	3-amino-2-methyl-3-phenylpropylamino
4-fluorophenyl	2-amino-4-pyrimidinyl	3-amino-2-methyl-3-phenylpropylamino
4-fluorophenyl	4-pyridyl	(S)-tetrahydroisoquinol-3-ylmethylenamino
4-fluorophenyl	2-acetamido-4-pyridyl	(S)-tetrahydroisoquinol-3-ylmethylenamino
4-fluorophenyl	2-amino-4-pyrimidinyl	(S)-tetrahydroisoquinol-3-ylmethylenamino
4-fluorophenyl	4-pyridyl	(S)-3-benzylpiperazinyl
4-fluorophenyl	2-acetamido-4-pyridyl	(S)-3-benzylpiperazinyl
4-fluorophenyl	2-amino-4-pyrimidinyl	(S)-3-benzylpiperazinyl

and

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in which R^2 is H, methyl or benzyl, and R^{11} , R^{12} , and R^1 are one of the combinations given in the following table:

R^{11}	R^{12}	R^1
Phenyl	4-pyridyl	phenyl
4-fluorophenyl	4-pyridyl	phenyl
Phenyl	2-acetamido-pyridyl	phenyl
4-fluorophenyl	2-acetamido-pyridyl	phenyl
Phenyl	4-pyridyl	4-ethylphenyl
4-fluorophenyl	4-pyridyl	4-ethylphenyl
Phenyl	2-acetamido-pyridyl	4-ethylphenyl
4-fluorophenyl	2-acetamido-pyridyl	4-ethylphenyl
Phenyl	4-pyridyl	2,4-dimethylphenyl
4-fluorophenyl	4-pyridyl	2,4-dimethylphenyl
Phenyl	2-acetamido-pyridyl	2,4-dimethylphenyl
4-fluorophenyl	2-acetamido-pyridyl	2,4-dimethylphenyl
Phenyl	4-pyridyl	2,6-dichlorobenzyl
4-fluorophenyl	4-pyridyl	2,6-dichlorobenzyl
Phenyl	2-acetamido-pyridyl	2,6-dichlorobenzyl
4-fluorophenyl	2-acetamido-pyridyl	2,6-dichlorobenzyl
Phenyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
4-fluorophenyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
Phenyl	2-acetamido-pyridyl	2-(4-fluorophenyl)ethylamino
4-fluorophenyl	2-acetamido-pyridyl	2-(4-fluorophenyl)ethylamino
Phenyl	4-pyridyl	3-phenylpropylamino
4-fluorophenyl	4-pyridyl	3-phenylpropylamino
Phenyl	2-acetamido-pyridyl	3-phenylpropylamino
4-fluorophenyl	2-acetamido-pyridyl	3-phenylpropylamino
Phenyl	4-pyridyl	1-piperazinyl
4-fluorophenyl	4-pyridyl	1-piperazinyl

Phenyl	2-acetamido-pyridyl	1-piperazinyl
4-fluorophenyl	2-acetamido-pyridyl	1-piperazinyl
benzyl	4-pyridyl	3-phenylpropylamino
benzyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
2-thienyl	4-pyridyl	3-phenylpropylamino
2-thienyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
cyclohexyl	4-pyridyl	3-phenylpropylamino
cyclohexyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
tert-butyl	4-pyridyl	3-phenylpropylamino
tert-butyl	4-pyridyl	2-(4-fluorophenyl)ethylamino
4-fluorophenyl	4-piperidinyl	3-phenylpropylamino
4-fluorophenyl	4-piperidinyl	2-(4-fluorophenyl)ethylamino
4-fluorophenyl	4-pyranyl	3-phenylpropylamino
4-fluorophenyl	4-pyranyl	2-(4-fluorophenyl)ethylamino
Phenyl	4-pyridyl	(S)-2-amino-3-phenylpropylamino
4-fluorophenyl	4-pyridyl	(S)-2-amino-3-phenylpropylamino
Phenyl	2-acetamido-pyridyl	(S)-2-amino-3-phenylpropylamino
4-fluorophenyl	2-acetamido-pyridyl	(S)-2-amino-3-phenylpropylamino
Phenyl	4-pyridyl	3-amino-3-phenylpropylamino
4-fluorophenyl	4-pyridyl	3-amino-3-phenylpropylamino
Phenyl	2-acetamido-pyridyl	3-amino-3-phenylpropylamino
4-fluorophenyl	2-acetamido-pyridyl	3-amino-3-phenylpropylamino
Phenyl	4-pyridyl	3-amino-2-methyl-3-phenylpropylamino
4-fluorophenyl	4-pyridyl	3-amino-2-methyl-3-phenylpropylamino
Phenyl	2-acetamido-pyridyl	3-amino-2-methyl-3-phenylpropylamino
4-fluorophenyl	2-acetamido-pyridyl	3-amino-2-methyl-3-phenylpropylamino
Phenyl	4-pyridyl	(S)-tetrahydroisoquinol-3-ylmethylenamino
4-fluorophenyl	4-pyridyl	(S)-tetrahydroisoquinol-3-ylmethylenamino
Phenyl	2-acetamido-pyridyl	(S)-tetrahydroisoquinol-3-ylmethylenamino

4-fluorophenyl	2-acetamido-pyridyl	(S)-tetrahydroisoquinol-3-ylmethylenamino
Phenyl	4-pyridyl	(S)-3-benzylpiperazinyl
4-fluorophenyl	4-pyridyl	S)-3-benzylpiperazinyl
Phenyl	2-acetamido-pyridyl	S)-3-benzylpiperazinyl
4-fluorophenyl	2-acetamido-pyridyl	S)-3-benzylpiperazinyl

Additional preferred compounds are listed in the Examples, *infra*.

As utilized herein, the following terms shall have
5 the following meanings:

"Alkyl", alone or in combination, means a straight-chain or branched-chain alkyl radical containing preferably 1-15 carbon atoms (C₁-C₁₅), more preferably 1-8 carbon atoms (C₁-C₈), even more preferably 1-6 carbon atoms (C₁-C₆), yet more preferably 1-4 carbon atoms (C₁-C₄), still more preferably 1-3 carbon atoms (C₁-C₃), and most preferably 1-2 carbon atoms (C₁-C₂). Examples of such radicals include methyl, ethyl, n-propyl, isopropyl,
10 n-butyl, isobutyl, sec-butyl, tert-butyl, pentyl, isoamyl, hexyl, octyl and the like.

"Hydroxyalkyl", alone or in combination, means an alkyl radical as defined above wherein at least one hydrogen radical is replaced with a hydroxyl radical, preferably 1-3 hydrogen radicals are replaced by hydroxyl radicals, more preferably 1-2 hydrogen radicals are replaced by hydroxyl radicals, and most preferably one hydrogen radical is replaced by a hydroxyl radical. Examples of such radicals include hydroxymethyl, 1-, 2-hydroxyethyl, 1-, 2-, 3-hydroxypropyl, 1,3-dihydroxy-2-propyl, 1,3-dihydroxybutyl, 1,2,3,4,5,6-hexahydroxy-2-hexyl and the like.
20

30 "Alkenyl", alone or in combination, means a straight-chain or branched-chain hydrocarbon radical having one

or more double bonds, preferably 1-2 double bonds and more preferably one double bond, and containing preferably 2-15 carbon atoms (C₂-C₁₅), more preferably 2-8 carbon atoms (C₂-C₈), even more preferably 2-6 5 carbon atoms (C₂-C₆), yet more preferably 2-4 carbon atoms (C₂-C₄), and still more preferably 2-3 carbon atoms (C₂-C₃). Examples of such alkenyl radicals include ethenyl, propenyl, 2-methylpropenyl, 1,4-butadienyl and the like.

10 "Alkoxy", alone or in combination, means a radical of the type "R-O-" wherein "R" is an alkyl radical as defined above and "O" is an oxygen atom. Examples of such alkoxy radicals include methoxy, ethoxy, n-propoxy, 15 isopropoxy, n-butoxy, iso-butoxy, sec-butoxy, tert-butoxy and the like.

20 "Alkoxycarbonyl", alone or in combination, means a radical of the type "R-O-C(O)-" wherein "R-O-" is an alkoxy radical as defined above and "C(O)" is a carbonyl radical.

25 "Alkoxycarbonylamino", alone or in combination, means a radical of the type "R-O-C(O)-NH-" wherein "R-O-C(O)" is an alkoxycarbonyl radical as defined above, wherein the amino radical may optionally be substituted, such as with alkyl, aryl, aralkyl, cycloalkyl, cycloalkylalkyl and the like.

30 "Alkylthio", alone or in combination, means a radical of the type "R-S-" wherein "R" is an alkyl radical as defined above and "S" is a sulfur atom. Examples of such alkylthio radicals include methylthio, ethylthio, n-propylthio, isopropylthio, n-butylthio, iso-butylthio, 35 sec-butylthio, tert-butylthio and the like.

"Alkylsulfinyl", alone or in combination, means a radical of the type "R-S(O)-" wherein "R" is an alkyl radical as defined above and "S(O)" is a mono-oxygenated sulfur atom. Examples of such alkylsulfinyl radicals 5 include methylsulfinyl, ethylsulfinyl, n-propylsulfinyl, isopropylsulfinyl, n-butylsulfinyl, iso-butylsulfinyl, sec-butylsulfinyl, tert-butylsulfinyl and the like.

"Alkylsulfonyl", alone or in combination, means a 10 radical of the type "R-S(O)₂-" wherein "R" is an alkyl radical as defined above and "S(O)₂" is a di-oxygenated sulfur atom. Examples of such alkylsulfonyl radicals include methylsulfonyl, ethylsulfonyl, n-propylsulfonyl, isopropylsulfonyl, n-butylsulfonyl, iso-butylsulfonyl, 15 sec-butylsulfonyl, tert-butylsulfonyl and the like.

"Aryl", alone or in combination, means a phenyl or biphenyl radical, which is optionally benzo fused or heterocyclo fused and which is optionally substituted 20 with one or more substituents selected from alkyl, alkoxy, halogen, hydroxy, amino, azido, nitro, cyano, haloalkyl, carboxy, alkoxy carbonyl, cycloalkyl, alkanoylamino, amido, amidino, alkoxy carbonylamino, N-alkylamidino, alkylamino, dialkylamino, aminoalkyl, 25 alkylaminoalkyl, dialkylaminoalkyl, N-alkylamido, N,N-dialkylamido, aralkoxy carbonylamino, alkylthio, alkylsulfinyl, alkylsulfonyl, oxo and the like. Examples of aryl radicals are phenyl, o-tolyl, 4-methoxyphenyl, 2-(tert-butoxy)phenyl, 3-methyl-4- 30 methoxyphenyl, 2-CF₃-phenyl, 2-fluorophenyl, 2-chlorophenyl, 3-nitrophenyl, 3-aminophenyl, 3-acetamidophenyl, 2-amino-3-(aminomethyl)phenyl, 6-methyl-3-acetamidophenyl, 6-methyl-2-aminophenyl, 6-methyl-2,3-diaminophenyl, 2-amino-3-methylphenyl, 4,6- 35 dimethyl-2-aminophenyl, 4-hydroxyphenyl, 3-methyl-4-hydroxyphenyl, 4-(2-methoxyphenyl)phenyl, 2-amino-1-naphthyl, 2-naphthyl, 3-amino-2-naphthyl, 1-methyl-3-

amino-2-naphthyl, 2,3-diamino-1-naphthyl, 4,8-dimethoxy-2-naphthyl and the like.

"Aralkyl" and "arylalkyl", alone or in combination,
5 means an alkyl radical as defined above in which at least one hydrogen atom, preferably 1-2, is replaced by an aryl radical as defined above, such as benzyl, 1-, 2-phenylethyl, dibenzylmethyl, hydroxyphenylmethyl, methylphenylmethyl, diphenylmethyl,
10 dichlorophenylmethyl, 4-methoxyphenylmethyl and the like.

"Aralkoxy", alone or in combination, means an alkoxy radical as defined above in which at least one hydrogen atom, preferably 1-2, is replaced by an aryl radical as defined above, such as benzyloxy, 1-, 2-phenylethoxy, dibenzylmethoxy, hydroxyphenylmethoxy, methylphenylmethoxy, dichlorophenylmethoxy, 4-methoxyphenylmethoxy and the like.
15

20 "Aralkoxycarbonyl", alone or in combination, means a radical of the type "R-O-C(O)-" wherein "R-O-" is an aralkoxy radical as defined above and "-C(O)-" is a carbonyl radical.

25 "Alkanoyl", alone or in combination, means a radical of the type "R-C(O)-" wherein "R" is an alkyl radical as defined above and "-C(O)-" is a carbonyl radical. Examples of such alkanoyl radicals include acetyl,
30 trifluoroacetyl, hydroxyacetyl, propionyl, butyryl, valeryl, 4-methylvaleryl, and the like.

"Alkanoylamino", alone or in combination, means a radical of the type "R-C(O)-NH-" wherein "R-C(O)-" is an
35 alkanoyl radical as defined above, wherein the amino radical may optionally be substituted, such as with

alkyl, aryl, aralkyl, cycloalkyl, cycloalkylalkyl and the like.

"Aminocarbonyl", alone or in combination, means an amino
5 substituted carbonyl (carbamoyl) radical, wherein the
amino radical may optionally be mono- or di-substituted,
such as with alkyl, aryl, aralkyl, cycloalkyl,
cycloalkylalkyl, alkanoyl, alkoxy carbonyl,
aralkoxy carbonyl and the like.

10

"Aminosulfonyl", alone or in combination, means an amino substituted sulfonyl radical.

"Benzo", alone or in combination, means the divalent
15 radical C₆H₄= derived from benzene. "Benzo fused" forms
a ring system in which benzene and a cycloalkyl or aryl
group have two carbons in common, for example
tetrahydronaphthylene and the like.

20 "Bicyclic" as used herein is intended to include both
fused ring systems, such as naphthyl and β -carbolinyl,
and substituted ring systems, such as biphenyl,
phenylpyridyl and diphenylpiperazinyl.

25 "Cycloalkyl", alone or in combination, means a saturated
or partially saturated, preferably one double bond,
monocyclic, bicyclic or tricyclic carbocyclic alkyl
radical, preferably monocyclic, containing preferably 5-
12 carbon atoms (C₅-C₁₂), more preferably 5-10 carbon
30 atoms (C₅-C₁₀), even more preferably 5-7 carbon atoms
(C₅-C₇), which is optionally benzo fused or heterocyclo
fused and which is optionally substituted as defined
herein with respect to the definition of aryl. Examples
of such cycloalkyl radicals include cyclopentyl,
35 cyclohexyl, dihydroxycyclohexyl,
ethylenedioxycyclohexyl, cycloheptyl, octahydronaphthyl,
tetrahydronaphthyl, octahydroquinolinyl,

dimethoxytetrahydronaphthyl, 2,3-dihydro-1H-indenyl, azabicyclo[3.2.1]octyl and the like.

"Heteroatoms" means nitrogen, oxygen and sulfur
5 heteroatoms.

"Heterocyclo fused" forms a ring system in which a heterocyclyl or heteroaryl group of 5-6 ring members and a cycloalkyl or aryl group have two carbons in common,
10 for example indole, isoquinoline, tetrahydroquinoline, methylenedioxybenzene and the like.

"Heterocyclyl" means a saturated or partially unsaturated, preferably one double bond, monocyclic or
15 bicyclic, preferably monocyclic, heterocycle radical containing at least one, preferably 1 to 4, more preferably 1 to 3, even more preferably 1-2, nitrogen, oxygen or sulfur atom ring member and having preferably 3-8 ring members in each ring, more preferably 5-8 ring
20 members in each ring and even more preferably 5-6 ring members in each ring. "Heterocyclyl" is intended to include sulfone and sulfoxide derivatives of sulfur ring members and N-oxides of tertiary nitrogen ring members, and carbocyclic fused, preferably 3-6 ring carbon atoms
25 and more preferably 5-6 ring carbon atoms, and benzo fused ring systems. "Heterocyclyl" radicals may optionally be substituted on at least one, preferably 1-4, more preferably 1-3, even more preferably 1-2, carbon atoms by halogen, alkyl, alkoxy, hydroxy, oxo, thioxo,
30 aryl, aralkyl, heteroaryl, heteroaralkyl, amidino, N-alkylamidino, alkoxy carbonylamino, alkylsulfonylamino and the like, and/or on a secondary nitrogen atom by hydroxy, alkyl, aralkoxycarbonyl, alkanoyl, alkoxy carbonyl, heteroaralkyl, aryl or aralkyl radicals.
35 More preferably, "heterocyclyl", alone or in combination, is a radical of a monocyclic or bicyclic saturated heterocyclic ring system having 5-8 ring

members per ring, wherein 1-3 ring members are oxygen, sulfur or nitrogen heteroatoms, which is optionally partially unsaturated or benzo-fused and optionally substituted by 1-2 oxo or thioxo radicals. Examples of such heterocyclyl radicals include pyrrolidinyl, piperidinyl, piperazinyl, morpholinyl, thiamorpholinyl, 4-benzyl-piperazin-1-yl, pyrimidinyl, tetrahydrofuryl, pyrazolidinyl, pyrazolinyl, pyridazinonyl, pyrrolidonyl, tetrahydrothienyl and its sulfoxide and sulfone derivatives, 2,3-dihydroindolyl, tetrahydroquinolinyl, 1,2,3,4-tetrahydroisoquinolinyl, 1,2,3,4-tetrahydro-1-oxo-isoquinolinyl, 2,3-dihydrobenzofuryl, benzopyranyl, methylenedioxyphenyl, ethylenedioxyphenyl and the like.

"Heteroaryl" means a monocyclic or bicyclic, preferably monocyclic, aromatic heterocycle radical, having at least one, preferably 1 to 4, more preferably 1 to 3, even more preferably 1-2, nitrogen, oxygen or sulfur atom ring members and having preferably 5-6 ring members in each ring, which is optionally saturated carbocyclic fused, preferably 3-4 carbon atoms (C₃-C₄) to form 5-6 ring membered rings and which is optionally substituted as defined above with respect to the definitions of aryl. Examples of such heteroaryl groups include imidazolyl, 1-benzyloxycarbonylimidazol-4-yl, pyrrolyl, pyrazolyl, pyridyl, 3-(2-methyl)pyridyl, 3-(4-trifluoromethyl)pyridyl, pyrimidinyl, 5-(4-trifluoromethyl)pyrimidinyl, pyrazinyl, triazolyl, furyl, thienyl, oxazolyl, thiazolyl, indolyl, quinolinyl, 5,6,7,8-tetrahydroquinolyl, 5,6,7,8-tetrahydroisoquinolinyl, quinoxalinyl, benzothiazolyl, benzofuryl, benzimidazolyl, benzoxazolyl and the like.

"Heteroaralkyl" and "heteroarylalkyl," alone or in combination, means an alkyl radical as defined above in which at least one hydrogen atom, preferably 1-2, is

replaced by a heteroaryl radical as defined above, such as 3-furylpropyl, 2-pyrrolyl propyl, chloroquinolinylmethyl, 2-thienylethyl, pyridylmethyl, 1-imidazolylethyl and the like.

5

"Halogen" and "halo", alone or in combination, means fluoro, chloro, bromo or iodo radicals.

"Haloalkyl", alone or in combination, means an alkyl 10 radical as defined above in which at least one hydrogen atom, preferably 1-3, is replaced by a halogen radical, more preferably fluoro or chloro radicals. Examples of such haloalkyl radicals include 1,1,1-trifluoroethyl, chloromethyl, 1-bromoethyl, fluoromethyl, 15 difluoromethyl, trifluoromethyl, bis(trifluoromethyl)methyl and the like.

"Pharmacologically acceptable salt" means a salt prepared by conventional means, and are well known by 20 those skilled in the art. The "pharmacologically acceptable salts" include basic salts of inorganic and organic acids, including but not limited to hydrochloric acid, hydrobromic acid, sulphuric acid, phosphoric acid, methanesulphonic acid, ethanesulfonic acid, malic acid, 25 acetic acid, oxalic acid, tartaric acid, citric acid, lactic acid, fumaric acid, succinic acid, maleic acid, salicylic acid, benzoic acid, phenylacetic acid, mandelic acid and the like. When compounds of the invention include an acidic function such as a carboxy 30 group, then suitable pharmaceutically acceptable cation pairs for the carboxy group are well known to those skilled in the art and include alkaline, alkaline earth, ammonium, quaternary ammonium cations and the like. For additional examples of "pharmacologically acceptable 35 salts," see *infra* and Berge et al, *J. Pharm. Sci.* **66**, 1 (1977).

"Cytokine" means a secreted protein that affects the functions of other cells, particularly as it relates to the modulation of interactions between cells of the immune system or cells involved in the inflammatory response. Examples of cytokines include but are not limited to interleukin 1 (IL-1), preferably IL-1 β , interleukin 6 (IL-6), interleukin 8 (IL-8) and TNF, preferably TNF- α (tumor necrosis factor- α).

5 "TNF, IL-1, IL-6, and/or IL-8 mediated disease or disease state" means all disease states wherein TNF, IL-1, IL-6, and/or IL-8 plays a role, either directly as TNF, IL-1, IL-6, and/or IL-8 itself, or by TNF, IL-1, IL-6, and/or IL-8 inducing another cytokine to be released. For example, a disease state in which IL-1 plays a major role, but in which the production of or action of IL-1 is a result of TNF, would be considered mediated by TNF.

10 "Leaving group" generally refers to groups readily displaceable by a nucleophile, such as an amine, a thiol or an alcohol nucleophile. Such leaving groups are well known in the art. Examples of such leaving groups include, but are not limited to, N-hydroxysuccinimide, N-hydroxybenzotriazole, halides, triflates, tosylates and the like. Preferred leaving groups are indicated herein where appropriate.

15 "Protecting group" generally refers to groups well known in the art which are used to prevent selected reactive groups, such as carboxy, amino, hydroxy, mercapto and the like, from undergoing undesired reactions, such as nucleophilic, electrophilic, oxidation, reduction and the like. Preferred protecting groups are indicated herein where appropriate. Examples of amino protecting groups include, but are not limited to, aralkyl, substituted aralkyl, cycloalkenylalkyl and substituted cycloalkenyl

alkyl, allyl, substituted allyl, acyl, alkoxycarbonyl, aralkoxycarbonyl, silyl and the like. Examples of aralkyl include, but are not limited to, benzyl, ortho-methylbenzyl, trityl and benzhydryl, which can be 5 optionally substituted with halogen, alkyl, alkoxy, hydroxy, nitro, acylamino, acyl and the like, and salts, such as phosphonium and ammonium salts. Examples of aryl groups include phenyl, naphthyl, indanyl, anthracenyl, 9-(9-phenylfluorenyl), phenanthrenyl, durenyl and the like.

10 Examples of cycloalkenylalkyl or substituted cycloalkylenylalkyl radicals, preferably have 6-10 carbon atoms, include, but are not limited to, cyclohexenyl methyl and the like. Suitable acyl, alkoxycarbonyl and aralkoxycarbonyl groups include benzyloxycarbonyl, t-butoxycarbonyl, iso-butoxycarbonyl, benzoyl, substituted benzoyl, butyryl, acetyl, tri-fluoroacetyl, tri-chloro acetyl, phthaloyl and the like. A mixture of protecting groups can be used to protect the same amino group, such 15 as a primary amino group can be protected by both an aralkyl group and an aralkoxycarbonyl group. Amino protecting groups can also form a heterocyclic ring with the nitrogen to which they are attached, for example, 1,2-bis(methylene)benzene, phthalimidyl, succinimidyl, maleimidyl and the like and where these heterocyclic groups can further include adjoining aryl and cycloalkyl 20 rings. In addition, the heterocyclic groups can be mono-, di- or tri-substituted, such as nitrophthalimidyl. Amino groups may also be protected against undesired reactions, such as oxidation, through the formation of an 25 addition salt, such as hydrochloride, toluenesulfonic acid, trifluoroacetic acid and the like. Many of the amino protecting groups are also suitable for protecting carboxy, hydroxy and mercapto groups. For example, aralkyl groups. Alkyl groups are also suitable groups 30 for protecting hydroxy and mercapto groups, such as tert-butyl.

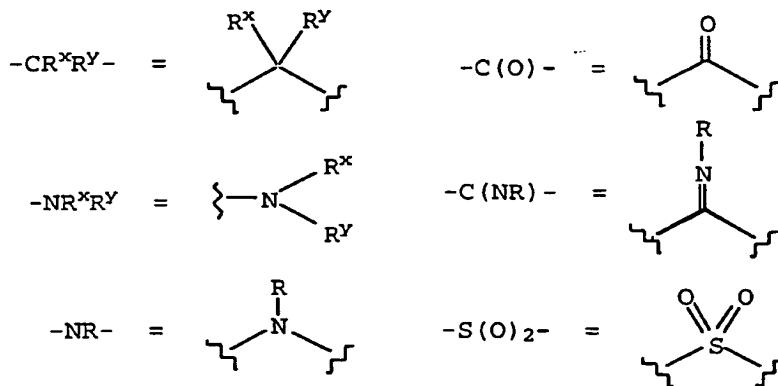
35

Silyl protecting groups are silicon atoms optionally substituted by one or more alkyl, aryl and aralkyl groups. Suitable silyl protecting groups include, but are not limited to, trimethylsilyl, 5 triethylsilyl, tri-isopropylsilyl, tert-butyldimethylsilyl, dimethylphenylsilyl, 1,2-bis(dimethylsilyl)benzene, 1,2-bis(dimethylsilyl)ethane and diphenylmethylsilyl. Silylation of an amino groups provide mono- or di-silylamino groups. Silylation of 10 aminoalcohol compounds can lead to a N,N,O-tri-silyl derivative. Removal of the silyl function from a silyl ether function is readily accomplished by treatment with, for example, a metal hydroxide or ammonium flouride reagent, either as a discrete reaction step or 15 in situ during a reaction with the alcohol group. Suitable silylating agents are, for example, trimethylsilyl chloride, tert-butyl-dimethylsilyl chloride, phenyldimethylsilyl chloride, diphenylmethyl silyl chloride or their combination products with 20 imidazole or DMF. Methods for silylation of amines and removal of silyl protecting groups are well known to those skilled in the art. Methods of preparation of these amine derivatives from corresponding amino acids, amino acid amides or amino acid esters are also well 25 known to those skilled in the art of organic chemistry including amino acid/amino acid ester or aminoalcohol chemistry.

Protecting groups are removed under conditions which will not affect the remaining portion of the 30 molecule. These methods are well known in the art and include acid hydrolysis, hydrogenolysis and the like. A preferred method involves removal of a protecting group, such as removal of a benzyloxycarbonyl group by 35 hydrogenolysis utilizing palladium on carbon in a suitable solvent system such as an alcohol, acetic acid, and the like or mixtures thereof. A t-butoxycarbonyl protecting group can be removed utilizing an inorganic

or organic acid, such as HCl or trifluoroacetic acid, in a suitable solvent system, such as dioxane or methylene chloride. The resulting amino salt can readily be neutralized to yield the free amine. Carboxy protecting group, such as methyl, ethyl, benzyl, tert-butyl, 4-methoxyphenylmethyl and the like, can be removed under hydrolysis and hydrogenolysis conditions well known to those skilled in the art.

The symbols used above have the following meanings:



10

Prodrugs of the compounds of this invention are also contemplated by this invention. A prodrug is an active or inactive compound that is modified chemically through in vivo physiological action, such as hydrolysis, metabolism and the like, into a compound of this invention following administration of the prodrug to a patient. The suitability and techniques involved in making and using prodrugs are well known by those skilled in the art. For a general discussion of prodrugs involving esters see Svensson and Tunek Drug Metabolism Reviews 165 (1988) and Bundgaard Design of Prodrugs, Elsevier (1985). Examples of a masked carboxylate anion include a variety of esters, such as alkyl (for example, methyl, ethyl), cycloalkyl (for example, cyclohexyl), aralkyl (for example, benzyl, p-methoxybenzyl), and alkylcarbonyloxyalkyl (for example, pivaloyloxymethyl). Amines have been masked as

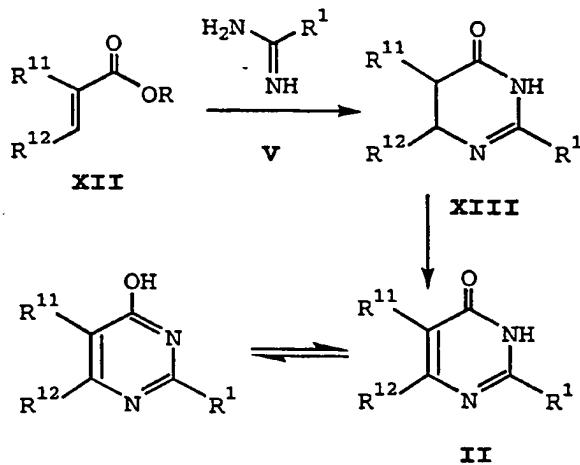
arylcarbonyloxymethyl substituted derivatives which are cleaved by esterases *in vivo* releasing the free drug and formaldehyde (Bungaard J. Med. Chem. 2503 (1989)). Also, drugs containing an acidic NH group, such as 5 imidazole, imide, indole and the like, have been masked with N-acyloxymethyl groups (Bundgaard Design of Prodrugs, Elsevier (1985)). Hydroxy groups have been masked as esters and ethers. EP 039,051 (Sloan and Little, 4/11/81) discloses Mannich-base hydroxamic acid 10 prodrugs, their preparation and use.

Compounds according to the invention can be synthesized according to one or more of the following methods. It should be noted that the general procedures are shown as it relates to preparation of compounds 15 having unspecified stereochemistry. However, such procedures are generally applicable to those compounds of a specific stereochemistry, e.g., where the stereochemistry about a group is (S) or (R). In addition, the compounds having one stereochemistry 20 (e.g., (R)) can often be utilized to produce those having opposite stereochemistry (i.e., (S)) using well-known methods, for example, by inversion.

4(3H)-Pyrimidinones:

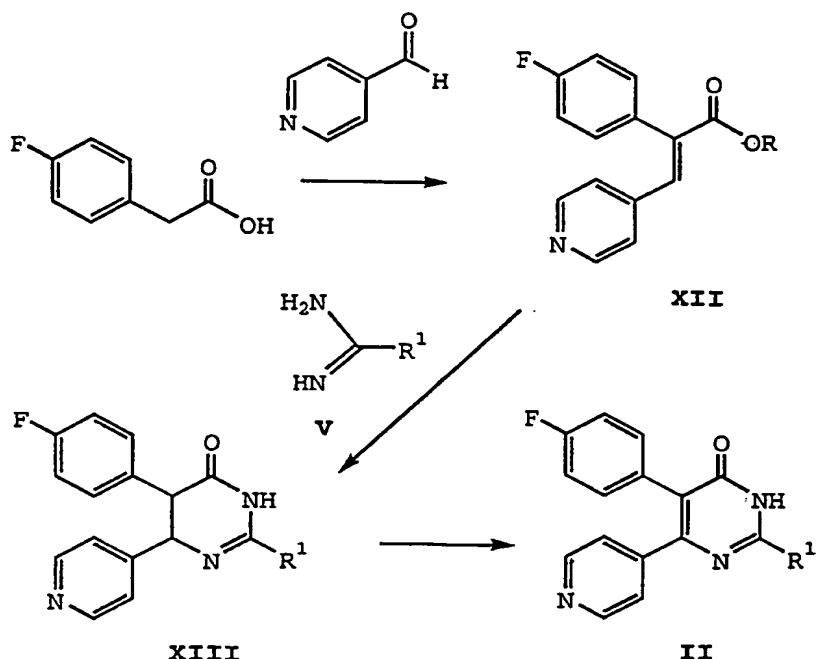
25 For the synthesis of 4(3H)-pyrimidinones II (or its tautomer, 4-hydroxy-pyrimidines), the approach displayed in Scheme 1 may be followed (for a review of synthetic methods see: D.J. Brown, Heterocyclic Compounds: the Pyrimidines, *supra*). This approach involves the 30 cyclization reaction between an acrylic acid ester XII and an amidine V followed by oxidation of the resulting dihydropyrimidinone XIII to give II.

Scheme 1



For the synthesis of 2-substituted 5-(4-fluorophenyl)-6-(4-pyridyl)-4-hydroxy-pyrimidines II (Scheme 2), the disubstituted acrylic acid ester XII may be prepared conveniently by condensation of pyridine-4-carboxaldehyde with 4-fluorophenylacetic acid followed by esterification. XII may be reacted with a variety of amidines V at elevated temperature. As a dehydrogenating agent for the conversion of XIII to II, sodium nitrite/acetic acid is suitable.

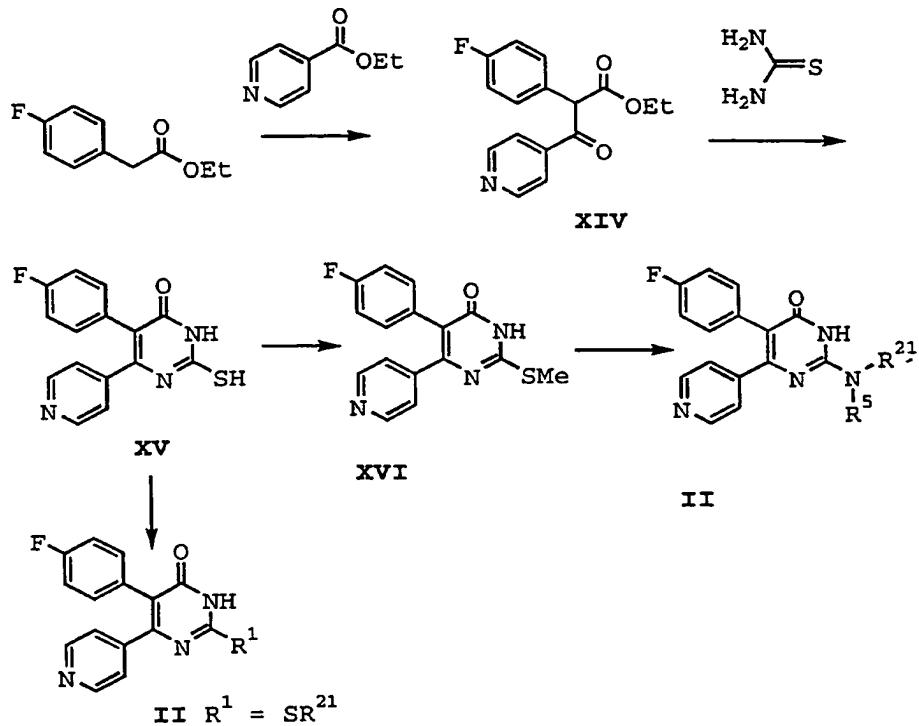
Scheme 2



Accordingly, further compounds of formula II may be obtained in which R^{12} is any other heteroaryl ring within the definition of R^{12} by the appropriate choice of starting material. Such starting materials include but are not limited to 2-methylpyridine-4-carboxaldehyde, 2,6-dimethylpyridine-4-carboxaldehyde (Mathes and Sauermilch, *Chem. Ber.* 88, 1276-1283 (1955)), quinoline-4-carboxaldehyde, pyrimidine-4-carboxaldehyde, 6-methylpyrimidine-4-carboxaldehyde, 2-methylpyrimidine-4-carboxaldehyde, 2,6-dimethylpyrimidine-4-carboxaldehyde (Bredereck et al., *Chem. Ber.* 97, 3407-3417 (1964)). The use of 2-nitropyridine-4-carboxaldehyde would lead to a derivative of formula II with R^{12} represented by a 2-nitro-4-pyridyl group. Catalytic reduction of the nitro to an amino group would provide the 2-amino-4-pyridyl derivative of II. The approach displayed in Scheme 2 is applicable to the use of other aryl acetic acids leading to compounds of formula II with different aryl groups as R^{11} .

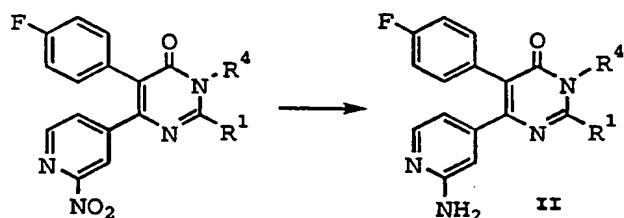
Pyrimidinone II ($R^1 = H$) may be substituted at the N-3 position by reaction with e.g. an alkyl halide, such as methyl iodide or ethyl bromide in the presence of an appropriate base such as potassium carbonate and the
5 like.

Scheme 3

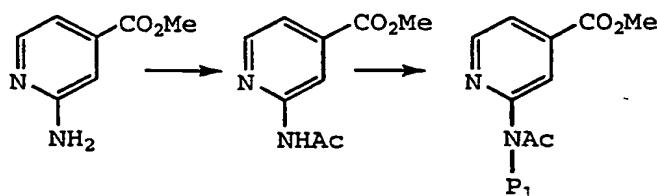


Another approach (Scheme 3) leading to 5,6-diaryl-4-hydroxy-pyrimidines involves the cyclization of the b-
10 keto ester XIV with thiourea to give the thiouracil derivative XV. XV can be S-monomethylated to XVI. Reaction of XVI with primary and secondary amines leads to 2-amino substituted 4-hydroxy-pyrimidines II. Further 2-thioether derivatives of II with $R^1 = SR^{21}$ can
15 be obtained, for example by alkylation of XV with alkyl halides. Treatment of XV or XVI with Raney nickel and H_2 provides compounds of structure II wherein R^1 is H.
Although Scheme 3 illustrates syntheses in which R^{12} is 4-pyridyl, this approach may be equally applied to

any other heteroaryl ring within the definition of R¹² by the appropriate choice of the starting material. Such starting materials include but are not limited to ethyl 2-methyl isonicotinate (Efimovsky and Rumpf, *Bull. Soc. Chim. FR.* 648-649 (1954)), methyl pyrimidine-4-carboxylate, methyl 2-methylpyrimidine-4-carboxylate, methyl 6-methylpyrimidine-4-carboxylate and methyl 2,6-dimethylpyrimidine-4-carboxylate (Sakasi et al., *Heterocycles* 13, 235 (1978)). Likewise, methyl 2-nitroisonicotinate (Stanonis, *J. Org. Chem.* 22, 475 (1957)) may be reacted with an aryl acetic acid ester followed by cyclization of the resultant β -keto ester with thiourea analogously to Scheme 3. Subsequent catalytic reduction of the nitro group to an amino group would give a pyrimidinone II in which R¹² is represented by a 2-amino-4-pyridyl group (Scheme 4).

Scheme 4

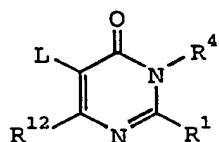
Furthermore, methyl 2-acetamido isonicotinate (Scheme 5) may be reacted analogously to Scheme 3 after appropriate protection of the amide nitrogen with e.g. a tert-butyldimethylsilyloxymethyl group (Benneche et al., *Acta Chem. Scand. B* 42 384-389 (1988)), a tert-butyldimethylsilyl group, a benzyloxymethyl group, a benzyl group or the like (P₁).

Scheme 5

Removal of the protecting group P₁ of the resulting pyrimidine II with a suitable reagent (e.g., tetrabutylammonium fluoride in the case where P₁ is t-butyldimethyl-silyloxymethyl) would then lead to a 5 pyrimidinone II with R¹² represented by a 2-acetamido-4-pyridyl group. Needless to say, ethyl p-fluorophenyl acetate may be substituted by any alkyl arylacetate in the procedure illustrated in Scheme 3 thus providing compounds of formula II with different R¹¹ aryl substituents.

10 In a further process, pyrimidinones II may be prepared by coupling a suitable derivative of XVIII (L is a leaving group, such as halogen radical and the like) with an appropriate aryl equivalent.

15



XVIII

Such aryl/heteroaryl couplings are well known to those skilled in the art and involve an organic-metallic component for reaction with a reactive derivative, e.g., 20 a halogeno derivative, of the second compound in the presence of a catalyst. The metallo-organic species may be provided either by the pyrimidinone in which case the aryl component provides the reactive halogen equivalent or the pyrimidinone may be in the form of a reactive 5-halogeno derivative for reaction with a metallo organic aryl compound. Accordingly, 5-bromo and 5-iodo derivatives of XVIII (L = Br, I) may be treated with arylalkyl tin compounds, e.g., trimethylstannylbenzene, in an inert solvent such as tetrahydrofuran in the 25 presence of a palladium catalyst, such as di(triphenylphosphine)palladium(II)dichloride. (Peters et al., *J. Heterocyclic Chem.* 27, 2165-2173, (1990). Alternatively, the halogen derivative of XVIII may be 30 converted into a trialkyltin derivative (L = Bu₃Sn) by

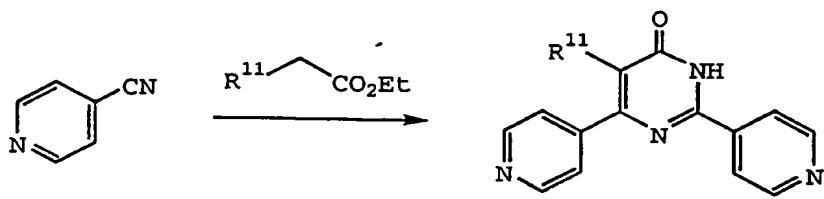
reaction with e.g. tributylstannyll chloride following lithiation with butyllithium and may then be reacted with an aryl halide in the presence of a catalyst.

(Sandosham and Undheim, *Acta Chem. Scand.* 43, 684-689

5 (1989)). Both approaches would lead to pyrimidines II in which R¹¹ is represented by aryl and heteroaryl groups.

As reported in the literature (Kabbe, *Lieb. Ann. Chem.* 704, 144 (1967); German Patent 1271116 (1968)) and displayed in Scheme 6, 5-aryl-2,6-dipyridyl-4(3H)-10 pyrimidinones II may be prepared in a one step synthesis by reaction of the cyanopyridine with an arylacetyl ester, such as ethyl phenylacetate in the presence of sodium methoxide.

Scheme 6



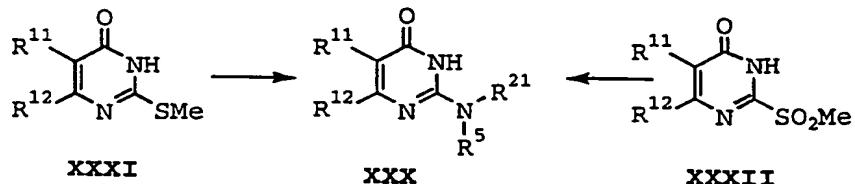
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In Scheme 7, compounds of the present invention of formula XXX can be readily prepared by reacting the methylthio intermediate XXXI with the amine NHR⁵R²¹, for example by heating the mixture preferably at a

20 temperature greater than 100°C, more preferably 150-210°C. Alternatively, compounds of formula XXX can be readily prepared by reacting the methylsulfonyl intermediate XXXII with the amine NHR⁵R²¹, for example by heating the mixture preferably at a temperature greater than 40°C, more preferably 50-210°C.

25

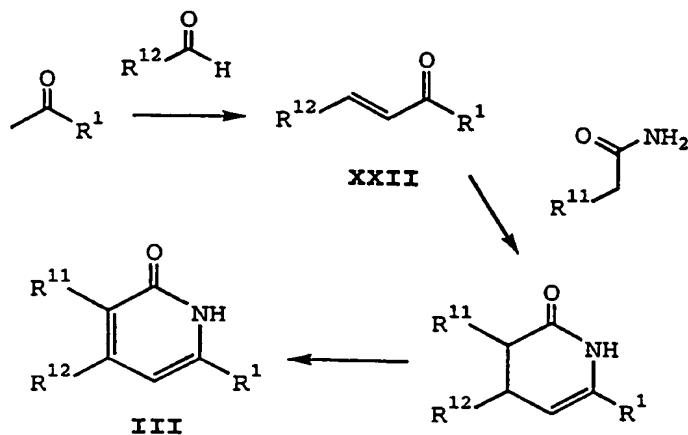
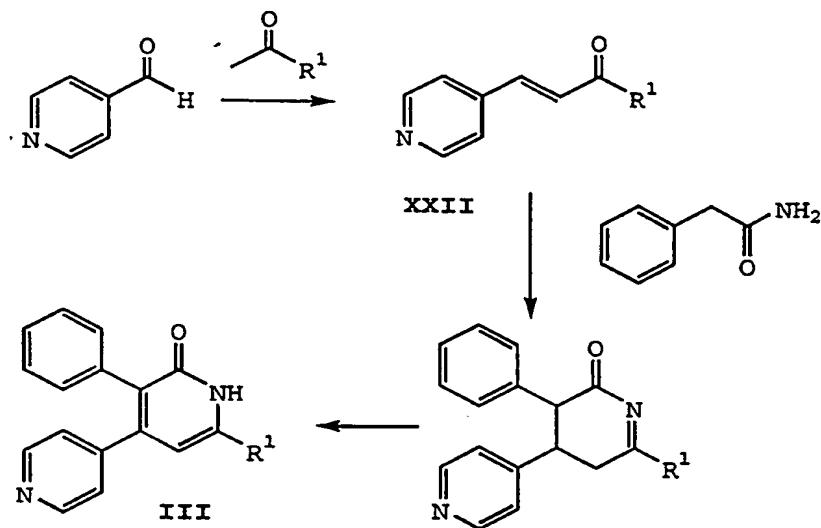
Scheme 7



Amines of formula $\text{NHR}'\text{R}''$ ²¹ are commercially available or can be readily prepared by those skilled in the art from commercially available starting materials. For example, an amide, nitro or cyano group can be reduced under reducing conditions, such as in the presence of a reducing agent like lithium aluminum hydride and the like, to form the corresponding amine. Alkylation and acylation of amino groups are well known in the art. Chiral and achiral substituted amines can be prepared from chiral amino acids and amino acid amides (for example, alkyl, aryl, heteroaryl, cycloalkyl, arylalkyl, heteroarylalkyl, cycloalkylalkyl and the like substituted glycine, β -alanine and the like) using methods well known in the art, such as H. Brunner, P. Hankofer, U. Holzinger, B. Treitinger and H. Schoenenberger, Eur. J. Med. Chem. 25, 35-44, 1990; M. Freiberger and R. B. Hasbrouck, J. Am. Chem. Soc. 82, 696-698, 1960; Dornow and Fust, Chem. Ber. 87, 984, 1954; M. Kojima and J. Fujita, Bull. Chem. Soc. Jpn. 55, 1454-1459, 1982; W. Wheeler and D. O'Bannon, Journal of Labelled Compounds and Radiopharmaceuticals XXXI, 306, 1992; and S. Davies, N. Garrido, O. Ichihara and I. Walters, J. Chem. Soc., Chem. Commun. 1153, 1993.

25 Pyridones:

As displayed in Scheme 8, a suitable route to 2(1*H*)-pyridones III involves the cyclization reaction between an *a,b*-unsaturated ketone XXII and a sufficiently reactive, substituted acetamide in the presence of base (El-Rayyes and Al-Hajjar, J. Heterocycl. Chem. 21, 1473 (1984)) and subsequent dehydrogenation.

Scheme 8Scheme 9

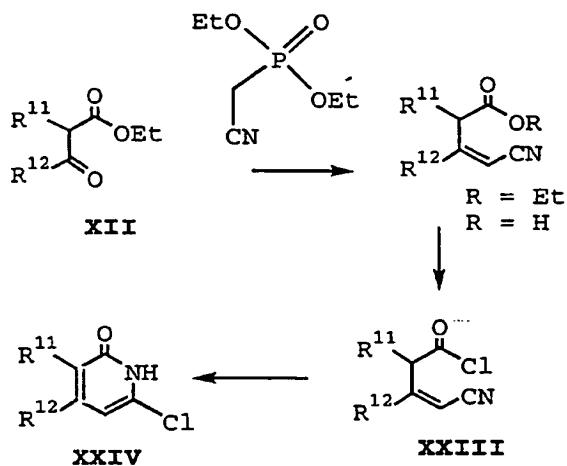
5 Accordingly (Scheme 9), pyridine-4-carboxaldehyde or other heteroaromatic carboxaldehyde-like pyrimidine-4-carboxaldehydes or quinoline-4-carboxyaldehydes may be reacted with acetyl aryl, acetyl heteroaryl or acetyl cycloalkyl derivatives in the presence of piperidine/acetic acid at elevated temperature (Bayer and Hartmann, *Arch. Pharm. (Weinheim)* 324, 815 (1991)) as well as pinacolone ($\text{CH}_3\text{-CO-C(CH}_3\text{)}_2$) in the presence of sodium hydroxide to provide the unsaturated ketone XXII (or the

10

analogous ketone from the corresponding heteroaromatic-4-carboxyaldehyde). The reaction of XXII with phenylacetamide in the presence of sodium ethoxide then may lead via the 3,4-dihydropyridone to 6-substituted 3-phenyl-4-(heteroaryl)-2(1H)-pyridones of structure III.

In Scheme 10, a feasible route is illustrated leading to 6-chloro-2(1H)-pyridone XXIV, a versatile intermediate for further modifications at the 6-position. This approach (G. Simchen, Chem. Ber. 103, 10 389-397 (1970) is based on the conversion of the unsaturated α -cyanocarboxylic acid chloride XXIII into XXIV in the presence of hydrogen chloride.

Scheme 10



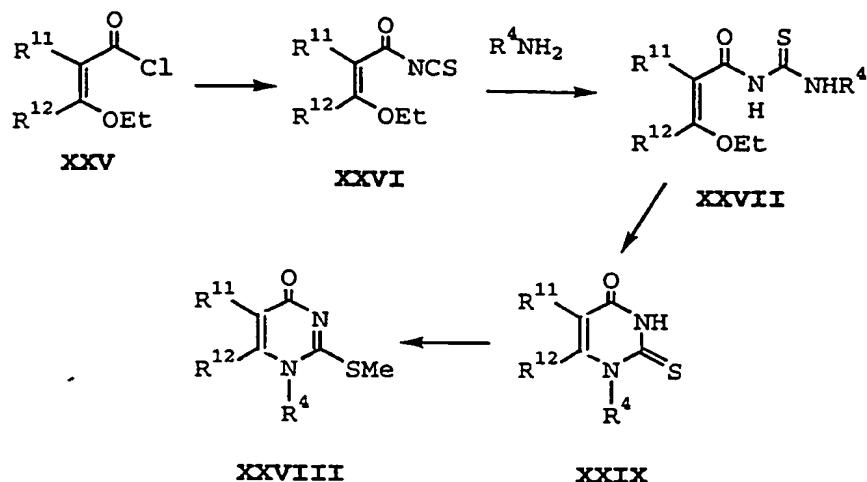
Reaction of XXIV with ammonia (Katritzky and Rachwal, J. Heterocyclic Chem. 32, 1007 (1995)), primary and secondary amines would lead to 2-amino substituted pyridones III. Furthermore, XXIV may be reacted in a palladium or nickel catalyzed cross-coupling reaction with an alkyl or aryl boronic acid or an alkyl or aryl zinc halide to provide pyridone III wherein R' is alkyl or aryl or heteroaryl.

In addition, pyridone III may be substituted at the N-1 position by reaction with, e.g., an alkyl halide in

the presence of an appropriate base such as potassium carbonate.

An approach that may lead to a pyrimidinone of the general formula III is illustrated in Scheme 11.

5

Scheme 11

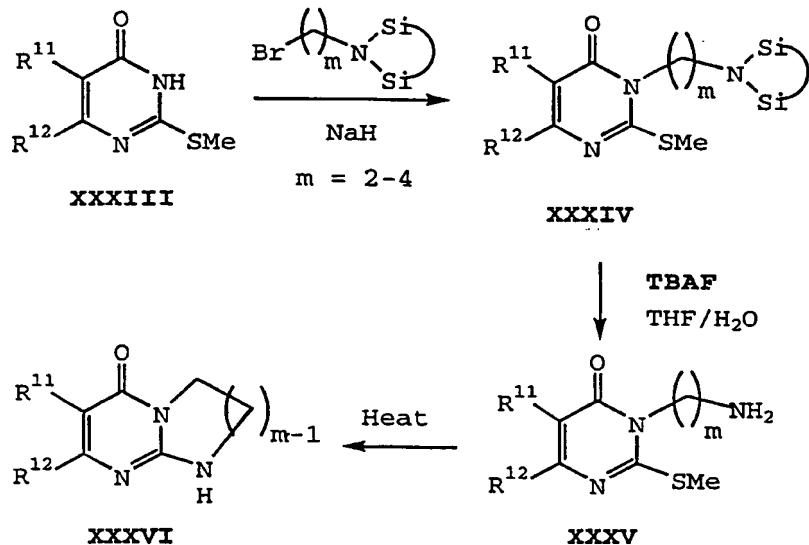
According to this approach (Shaw and Warrener, *J. Chem. Soc.* 153-156 (1958); Hronowski and Szarek, *Can. J. Chem.* 63, 2787 (1985); Agathocleous and Shaw, *J. Chem. Soc. Perkin Trans. I*, 2555 (1993)), an ethoxyacryloyl isothiocyanate XXVI is reacted with a primary amine to give as an addition product the acylthiourea XXVII which can be cyclized under basic or acidic conditions to the thiouracil compound XXVIII. XXVIII may be methylated to the methylthio derivative XXIX, a versatile intermediate for further transformations at the 2-position.

Fused 4(3*H*)-Pyrimidinones:

As displayed in Schemes 12 and 13, introduction of a suitable R' group through the alkylation of XXXIII affords an intermediate to the fused 5, 6, or 7 membered ring systems of Formula I wherein R' and V or W are joined. The synthesis utilizes a haloalkylamine in which the amino group is protected through reaction with 1,2-bis(chlorodimethylsilyl)ethane affording the cyclic

stabase derivative (see: Basha and Debernardis Tetrahedron Lett 5271, 1984) which protects the amine in the subsequent alkylation step (sodium hydride, DMF).

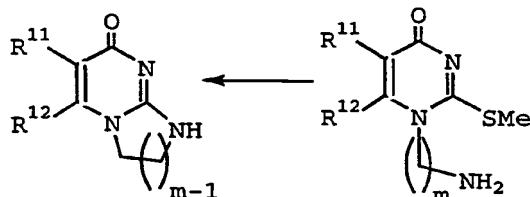
Scheme 12



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XXXVI

Scheme 13

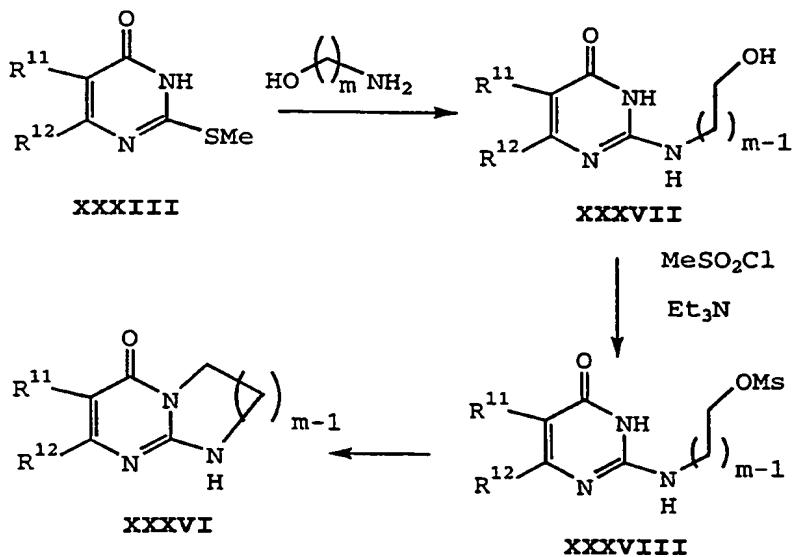


Deprotection of the amine can be accomplished with acid treatment (p-toluenesulfonic acid) or tetrabutylammonium fluoride treatment. The free amine can then be cyclized in an intramolecular fashion by warming to high temperatures. The bromoalkylamines are either commercially available (eg. 3-bromopropylamine hydrobromide, 2-bromoethylamine hydrobromide) or they can be synthesized from the corresponding haloalkylazide followed by reduction of the azide to the amine (see: Hendry et al Tetrahedron Lett 4597 (1987)). More functionalized haloalkylamines can be used as long as the functional groups are tolerated in the

transformations shown in scheme 12 including the bromo derivatives obtained from amino acid precursors as described by Baldwin et al (Synlett. 51-53, 1993) and Leanna et al (Tetrahedron Lett. 4485, 1993).

5 Alternatively, the fused ring system can be made through the addition of a hydroxyalkylamine as outlined in Scheme 14. Initially, the amine component of the hydroxyalkylamine displaces the 2-methylthio group to afford compound XXXVII which is followed by conversion
10 of the alcohol to a suitable leaving group (eg. methanesulfonate or trifluoromethanesulfonate). Closure of the ring can be accomplished by treatment with an excess of sodium hydride in DMF to afford XXXVI.

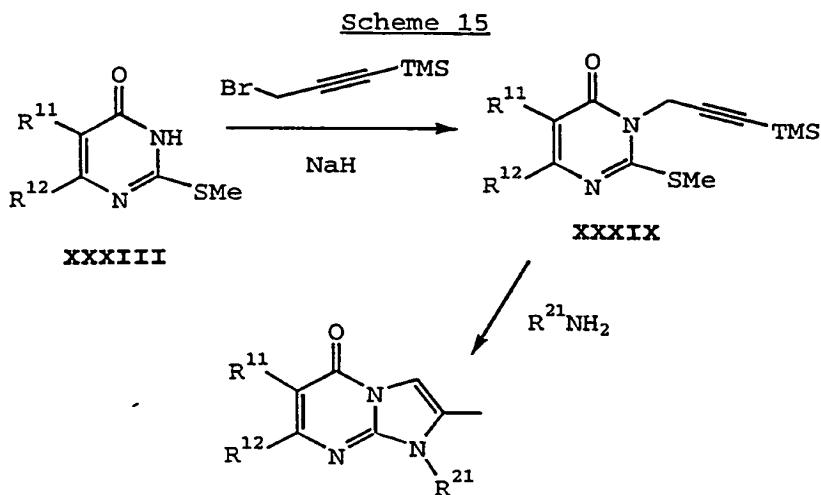
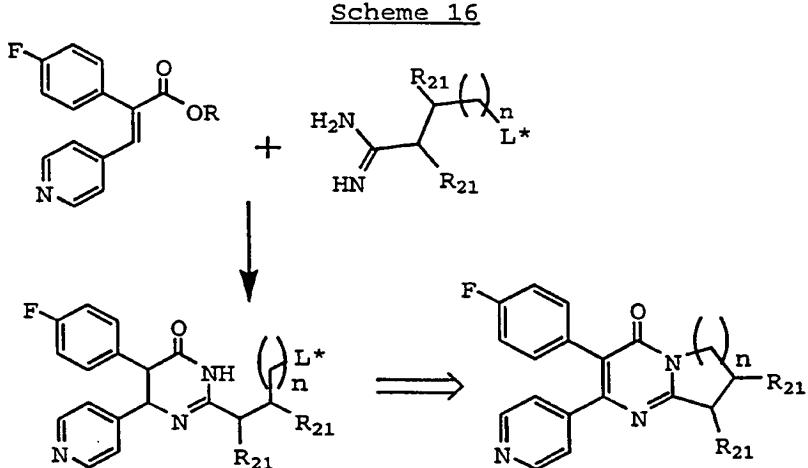
Scheme 14



15 The 6,5 fused ring systems can be obtained as outlined in Scheme 15. Alkylation of the N-3 nitrogen with 3-bromo-1-trimethylsilylpropyne can be followed by a displacement of the 2-methylthio group with the appropriate amine component exemplified but not limited to a phenylalkylamine. The 2-amino group under the reaction conditions cyclizes onto the acetylene as shown with a loss of the trimethylsilyl group as well. This transformation is illustrated in the examples below

wherein 3-phenyl-1-propylamine and benzylamine are reacted with 3-(3-trimethylsilyl-2-propynyl)-5-(4-fluorophenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone to afford the corresponding 6, 5 fused system.

5

Scheme 16

10

Compounds of the invention when U is CHR_{21} , can be prepared according to Scheme 2 above wherein R1 contains an leaving group or a group which can be converted into a leaving group (L^*) which can be reacted with a

primidine nitrogen atoms to form the fused ring (see Scheme 16).

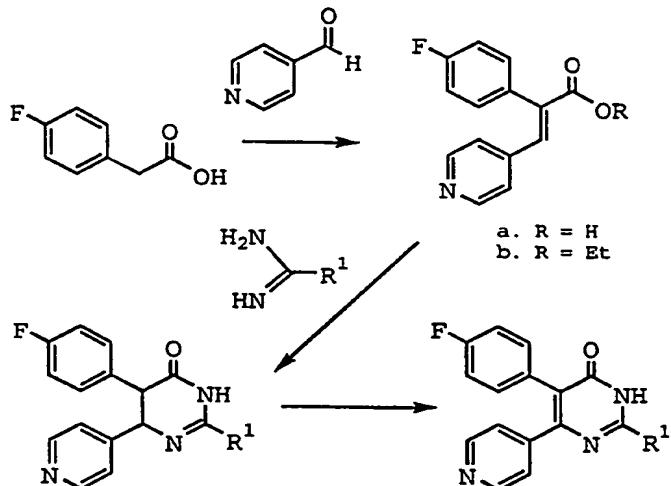
The following Examples are presented for illustrative purposes only and are not intended, nor
5 should they be construed, as limiting the invention in any manner. Those skilled in the art will appreciate that modifications and variations of the compounds disclosed herein can be made without violating the spirit or scope of the present invention.

10

EXAMPLES

Example 1

General procedure for the preparation of 2-substituted 5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidones



15 a. 2-(4-Fluorophenyl)-3-(4-pyridyl)-acrylic acid: A mixture of 4-fluorophenylacetic acid (9 g, 58.4 mmol), 4-pyridinecarboxaldehyde (5.6 ml, 58.6 mmol), pyridine (6 ml) and acetic anhydride (6 ml) was heated at 150°C for 1 h followed by evaporation and co-distillation with water. The resulting material crystallized on addition of ethanol. The solids were filtered and washed with ethanol and ethyl acetate to provide the title compound.
20 MS (m/z): 244.0 (M+H)⁺; C₁₄H₁₀FNO₂ requir. 243.2 ¹H-NMR

(DMSO-d₆): d 8.43, 6.98 (2d, each 2H, Pyrid.), 7.73 (s, 1H, CH=), 7.21 (d, 4H, PhF).

b. Ethyl 2-(4-fluorophenyl)-3-(4-pyridyl)-acrylate:

Conc. sulfuric acid (2.2 ml) was added carefully to a
5 suspension of 2-(4-fluorophenyl)-3-(4-pyridyl)-acrylic
acid (6.7 g, 27.5 mmol) in ethanol (120 ml) and the
mixture was heated at reflux for 24 h. The solvent was
evaporated, the remainder was taken up in
10 dichloromethane and the organic solution was washed with
aqueous sodium hydrogencarbonate and water, followed by
drying and evaporation. Flash column chromatography on
silica gel (hexane-acetone = 2:1) provided the pure
title compound. MS (m/z): 271.8 (M+H)⁺; C₁₆H₁₄FNO, requir.
271.3 ¹H-NMR (CDCl₃): 8.44, 6.88 (2m, each 2H, Pyrid.),
15 7.72 (s, 1H, CH=), 7.16, 7.06 (2m, each 2H, PhF), 4.28
(q, 2H, CH₂), 1.28 (t, 3H, CH₃).

c. General procedure: A stirred mixture of ethyl 2-(4-fluorophenyl)-3-(4-pyridyl)-acrylate (357 mg, 1.38 mmol), the amidine hydrochloride (2.61 mmol) and sodium
20 methoxide (250 mg, 4.62 mmol) in ethanol (5 ml) was
heated in a sealed tube at 120°C for 3 h. It was
neutralized with 2N hydrochloric acid prior to
evaporation. The residue was taken up in acetic acid
(25 ml) and treated with sodium nitrite (670 mg, 9.71
25 mmol) at 44°C for 20 min. After evaporation, the
resultant product was taken up in dichloromethane and
the solution was washed with aqueous sodium
hydrogencarbonate and water before drying and
evaporation. The product was purified by
30 recrystallization from methanol. If the crude product
of nitrite oxidation was water soluble, as was found for
5-(4-fluorophenyl)-2-methyl-6-(4-pyridyl)-4(3H)-
pyrimidinone, then no aqueous work up was done, but the
material obtained on evaporation was applied to a column
35 of silica gel (5% methanol/dichloromethane) prior to
recrystallization.

The following compounds were prepared accordingly using the appropriate amidine hydrochloride:

1-1 5-(4-Fluorophenyl)-2-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (m/z): 282.2 (M+H)⁺; C₁₆H₁₂FN₂O requir.

5 281.3 ¹H-NMR (DMSO-d₆): d 8.46 (m 2H, Pyrid.), 7.2-7.03 (m, 6H, PhF, Pyrid.). 2.38 (s, 3H, CH₃).

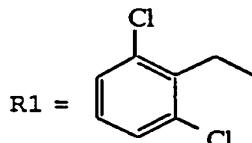
R1 = CH₃-

1-2 5-(4-Fluorophenyl)-2-isopropyl-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (m/z): 310.0 (M+H)⁺; C₁₈H₁₄FN₂O requir.

10 309.4 ¹H-NMR (DMSO-d₆): 8.45 (m, 2H, Pyrid.), 7.21-7.03 (m, 6H, PhF, Pyrid.), 2.90 (m, 1H, CH(CH₃)₂), 1.26, 1.24 (2s, each 3H, 2CH₃).

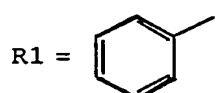
R1 = (CH₃)₂CH-

1-3 2-(2,6-Dichlorobenzyl)-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (m/z): 426.0 (M)⁺; C₂₂H₁₄Cl₂FN₂O requir. 426.3 ¹H-NMR (DMSO-d₆): d 8.37 (m, 2H, Pyrid.), 7.50 (d, 2H, PhCl₂), 7.35 (t, 1H, PhCl₂), 7.18-7.08 (m, 4H, PhF), 6.96 (m, 2H, Pyrid.), 4.36 (s, 2H, CH₂).



1-4 5-(4-Fluorophenyl)-2-phenyl-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (m/z): 344.2 (M+H)⁺; C₂₁H₁₄FN₂O requir.

25 343.4 ¹H-NMR (DMSO-d₆): d 8.49 (d, 2H, Pyrid.), 8.20 (d, 2H, Ph), 7.66-7.50 (m, 3H, Pyrid., Ph), 7.32-7.11 (m, 6H, PhF, Ph).

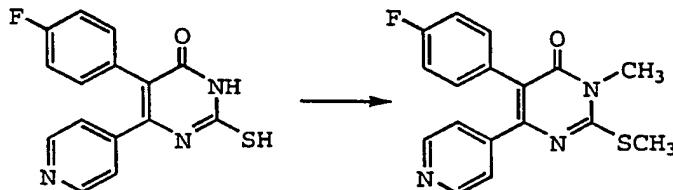


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Example 2

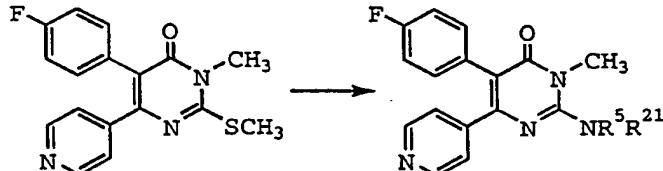
General procedure for the preparation of 2-N substituted 2-amino-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinones

5 **Step A. 5-(4-Fluorophenyl)-3-methyl-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone:**



Methyl iodide (418 ml, 6.67 mmol) was added to a stirred mixture of 5-(4-fluorophenyl)-6-(4-pyridyl)-2-thiouracil (1.0 g, 3.34 mmol) and potassium carbonate (923 mg, 6.68 mmol) in *N,N*-dimethylformamide (30 ml) at room temperature. Stirring was continued for 3 h, followed by evaporation and flash chromatography on a column of silica gel (hexane-acetone = 3:1, 2:1, 1:1) or 10 Iatrobeads[®] (chloroform-methanol = 90:7; chloroform-methanol-triethylamine = 90:7:3). The second main fraction provided the title compound as a solid. MS (m/z): 328.0 (M+H)⁺; C₁₁H₁₄FN₃OS requir. 327.4. ¹H-NMR (DMSO-d₆): δ 8.50, 7.26 (2m, each 2H, Pyrid.), 7.18, 15 7.14 (2m, each 2H, PhF), 3.52 (s, 3H, NCH₃), 2.65 (s, 3H, SCH₃).
20

Step B. General procedure:

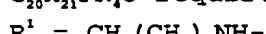


A mixture of 5-(4-fluorophenyl)-3-methyl-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone (103 mg, 0.32 mmol) and the amine HNR⁵R²¹ (1.2-3.2 mmol) was heated at 190-200°C for 2-48 h. The resulting product
25

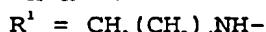
was purified by flash chromatography on a column of silica gel (hexane-acetone or methanol-dichloromethane or methanol-dichloromethane-conc. ammonium hydroxide) to provide the target compound.

5 The following compounds were prepared using the above procedure outlined above and an appropriate amine:
2-1 2-(n-Butylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone:

The reaction was done in a sealed tube at 190°C for 5 h.
10 MS (m/z): 353.0 (M+H)⁺; C₂₀H₂₁FN₄O requir. 352.4.

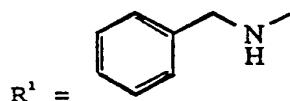


2-2 5-(4-Fluorophenyl)-3-methyl-2-(pentylamino)-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done in a
15 sealed tube at 190°C for 2.5 h. MS (m/z): 366.8 (M+H)⁺; C₂₁H₂₂FN₄O requir. 366.4.

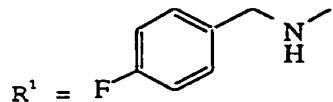


2-3 2-(3,3-Dimethylbutylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction
20 was done in a sealed tube at 190°C for 5 h. MS (m/z): 381.2 (M+H)⁺; C₂₂H₂₅FN₄O requir. 380.5.
R¹ = (CH₃)₂C(CH₂)₂NH-

2-4 2-(Benzylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at
25 185°C for 6h. MS (m/z): 387.2 (M+H)⁺; C₂₂H₂₁FN₄O requir. 386.4

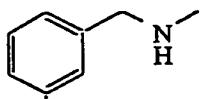


2-5 2-(4-Fluorobenzylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction
30 was done at 190°C for 24 h. MS (m/z): 405.2 (M+H)⁺; C₂₃H₁₈FN₄O requir. 404.4.

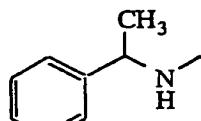


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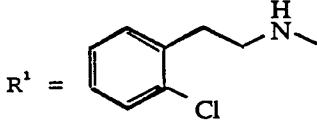
2-6 2-(3-Fluorobenzylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 195°C for 40 h. MS (m/z) : 405.0 (M+H)⁺; C₂₃H₁₈F₂N₄O requir. 404.4.

5 R¹ = F

2-7 5-(4-Fluorophenyl)-3-methyl-((R-1-phenylethyl)amino)-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 180°C for 4 days. MS (m/z) : 401.0 (M+H)⁺; C₂₄H₂₁FN₄O requir. 400.5

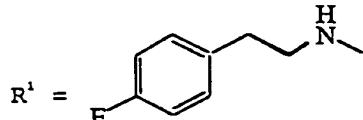
10 R¹ = F

2-8 2-(2-(2-Chlorophenyl)-ethylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 190°C for 5 h. MS (m/z) : 435.2 (M+H)⁺; C₂₄H₂₀ClFN₄O requir. 434.9.



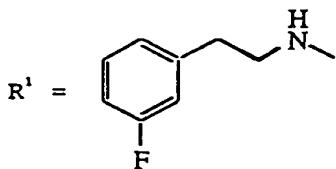
15

2-9 5-(4-Fluorophenyl)-2-(2-(4-fluorophenyl)-ethylamino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 190°C for 5 h. MS (m/z) : 419.2 (M+H)⁺; C₂₄H₂₀F₂N₄O requir. 418.5



20

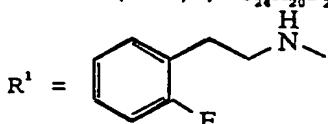
2-10 5-(2-Fluorophenyl)-2-(2-(3-fluorophenyl)-ethylamino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 190°C for 24 h. MS (m/z) : 419.2 (M+H)⁺; C₂₄H₂₀F₂N₄O requir. 418.5



2-11 5-(2-Fluorophenyl)-2-(2-(2-fluorophenyl)-ethylamino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone:

The reaction was done at 190°C for 12 h. MS (m/z):

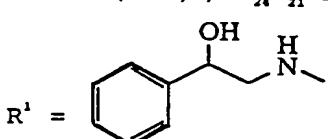
5 419.0 ($M+H$)⁺; $C_{24}H_{20}FN_4O$ requir. 418.5



2-12 5-(2-Fluorophenyl)-2-((2-hydroxy-2-phenyl)-ethylamino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone:

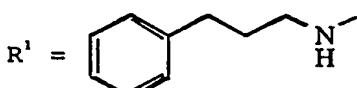
The reaction was done at 190°C for 1.5 h. MS (m/z):

10 417.0 ($M+H$)⁺; $C_{24}H_{21}FN_4O_2$ requir. 416.5.



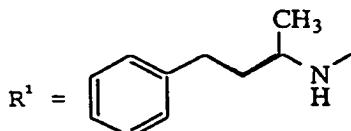
2-13 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 190°C for 6 h. MS (m/z): 415.0 ($M+H$)⁺;

15 $C_{22}H_{23}FN_4O$ requir. 414.5. $^1\text{H-NMR}$ (CDCl_3): d 8.49, 7.20 (2m, each 2H, Pyrid.), 7.35 (t, 2H, Ph), 7.30-7.25 (m, 3H, Ph), 7.12, 6.97 (2m, each 2H, PhF), 4.61 (t, 1H, NH), 3.67 (q, 2H, CH_2N), 3.28 (s, 3H, CH_3), 2.82 (t, 2H, CH_2Ph), 2.12 (m, 2H, CH_2).



20 2-14 5-(4-Fluorophenyl)-3-methyl-2-((1-methyl-3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone:

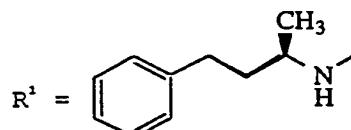
The reaction was done at 200°C for 48h. MS (m/z): 429.0 ($M+H$)⁺; $C_{26}H_{25}FN_4O$ requir. 428.5.



2-15 5-(4-Fluorophenyl)-3-methyl-2-((R)-1-methyl-3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone:

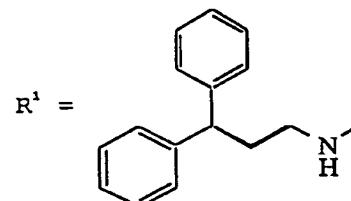
The reaction was done at 200°C for 48 h. MS (m/z) :

5 429.0 ($M+H$)⁺; $C_{26}H_{23}FN_4O$ requir. 428.5.



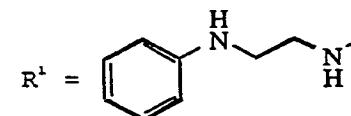
2-16 2-((3,3-Diphenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 190°C for 6 h. MS (m/z) : 490.8 ($M+H$)⁺;

10 $C_{31}H_{22}FN_4O$ requir. 490.6.



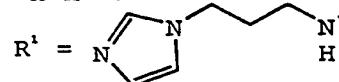
2-17 5-(4-Fluorophenyl)-3-methyl-2-((2-phenylaminoethyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 190°C for 4 h.

15 MS (m/z) : 416.2 ($M+H$)⁺; $C_{24}H_{22}FN_4O$ requir. 415.5.



2-18 5-(4-Fluorophenyl)-2-((3-imidazolylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 190°C for 2 h. MS (m/z) : 405.0 ($M+H$)⁺;

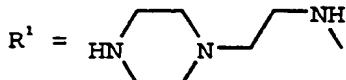
20 $C_{22}H_{21}FN_4O$ requir. 404.5.



2-19 5-(4-Fluorophenyl)-3-methyl-2-(2-(piperazin-1-yl)-ethylamino)-6-(4-pyridyl)-4(3H)-pyrimidinone: The

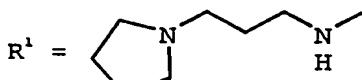
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reaction was done at 190°C for 30 min. MS (*m/z*): 409.2 (M+H)⁺; C₂₂H₂₄FN₆O requir. 408.5.



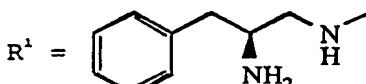
2-20 5-(4-Fluorophenyl)-3-methyl-6-(4-pyridyl)-2-(3-

5 (pyrrolidin-1-yl)-propylamino)-4(3H)-pyrimidinone: The reaction was done at 190°C for 2 h. MS (*m/z*): 408.2 (M+H)⁺; C₂₃H₂₆FN₆O requir. 407.5.



2-21 2-(((S)-2-Amino-3-phenylpropyl)-amino)-5-(4-

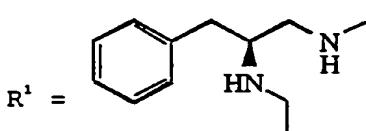
10 fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: The reaction was done at 190°C for 2.5 h. MS (*m/z*): 430.1 (M+H)⁺; C₂₅H₂₄FN₅O requir. 429.5 (free base).



15 2-22 2-(((S)-2-N-Ethyl-3-phenylpropyl)-amino)-5-(4-

fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: The reaction was done at 190°C for 4 h.

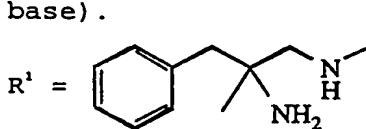
MS (*m/z*): 458.3 (M+H)⁺; C₂₇H₂₈FN₅O requir. 457.6 (free base).



20 2-23 2-((2-Amino-2-methyl-3-phenylpropyl) amino)-5-(4-

fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: The reaction was done at 190°C for 4 h.

MS (*m/z*): 444.0 (M+H)⁺; C₂₆H₂₆FN₅O requir. 443.5 (free base).

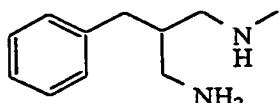


2-24 2-((2-Aminomethyl-3-phenylpropyl)-amino)-5-(4-

fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone

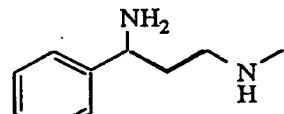
hydrochloride: The reaction was done at 190°C for 1 h.

MS (*m/z*): 444.0 (*M+H*)⁺; C₂₆H₂₆FN₄O requir. 443.5 (free base).



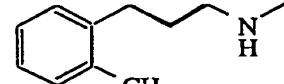
5 R¹ = 2-25 2-((3-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone

hydrochloride: The reaction was done at 190°C for 2.5 h. MS (*m/z*): 430.0 (*M+H*)⁺; C₂₅H₂₄FN₅O requir. 429.5 (free base).

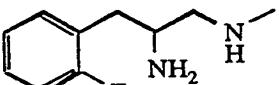


10 R¹ = 2-26 5-(4-Fluorophenyl)-3-methyl-2-(3-(2-methylphenyl)propyl)-amino-6-(4-pyridyl)-4(3H)-pyrimidinone

hydrochloride: The reaction was done at 190°C for 4 h. MS (*m/z*): 429.5 (*M+H*)⁺; C₂₆H₂₅FN₄O requir. 428.5.

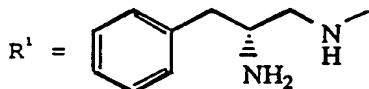


15 R¹ = 2-27 5-(4-Fluorophenyl)-3-methyl-2-((R,S)-2-amino-3-(2-fluorophenyl)-propyl-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone Hydrochloride: The reaction was done at 190°C for 7 h. MS (*m/z*): 448 (*M+H*)⁺.



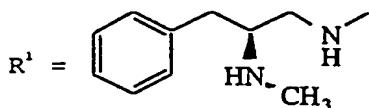
20 R¹ = 2-28 2-((R)-2-Amino-3-phenylpropyl)-amino-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone

hydrochloride: The reaction was done at 190°C for 2 h. MS (*m/z*): 430.2 (*M+H*)⁺; C₂₅H₂₄FN₅O requir. 429.5 (free base).



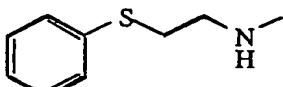
2-29 2-((S)-2-N-Methyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: The reaction was done at 190°C for 4 h.

5 MS (m/z): 444.0 (M+H)⁺; C₂₆H₂₆FN₅O requir. 443.5 (free base).

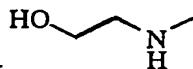


2-30 2-((2-phenylthioethyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction

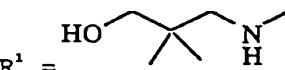
10 was done at 190°C for 16 h. MS (m/z): 433 (M+H)⁺.



$R^1 =$
2-31 2-((2-hydroxyethyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 190°C for 16 h. MS (m/z): 341 (M+H)⁺.

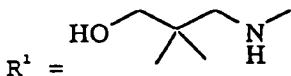


15 $R^1 =$
2-32 2-((2,2-dimethyl-3-hydroxypropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 190°C for 16 h. MS (m/z): 383 (M+H)⁺.



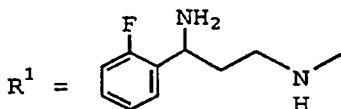
20 $R^1 =$
2-33 2-((2,2-dimethyl-3-phenylthiopropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: To a solution of triphenylphosphine (262 mg, 0.29 mmol) in tetrahydrofuran (2 mL) at 0°C was added diisopropyl 25 azodicarboxylate (DIAD) (56 mL, 0.29 mmol). After 30 min at 0°C, 2-((2,2-dimethyl-3-hydroxypropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone (50 mg, 0.14 mmol) and 2,6-dichlorothiophenol in tetrahydrofuran (2 mL) was added.

After 16 h, the reaction was concentrated under a stream of nitrogen. The reaction mixture was applied directly to purification via flash chromatography (step gradient ethyl acetate:CHCl₃ 1:3 then 1:2 then 1:1 then 2:1 then 5 3:1) to afford the title compound: MS (m/z) 544 (M+H)⁺.



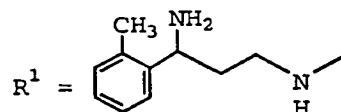
2-34 2-((3-Amino-3-(2-fluorophenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone

was prepared from 5-(4-fluorophenyl)-3-methyl-2-10 methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone and 1-(2-fluorophenyl)-1,3-propanediamine according to the General Procedure. The reaction was done at 190°C for 3 h. MS (m/z): 448.1 (M+H)⁺; C₂₅H₂₃F₂N₅O requir. 447.5 (free base).



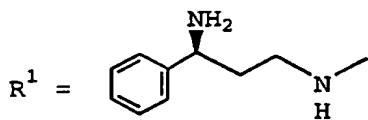
2-35 2-((3-Amino-3-(2-methylphenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone

hydrochloride was prepared from 5-(4-fluorophenyl)-3-methyl-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone 20 and 1-(2-methylphenyl)-1,3-propanediamine according to the General Procedure. The reaction was done at 185°C for 4 h. MS (m/z): 444.5 (M+H)⁺; C₂₆H₂₆FN₅O requir. 443.5 (free base).



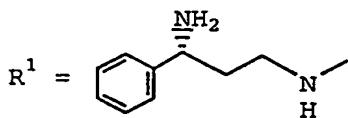
2-36 2-(((S)-3-amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone

hydrochloride was prepared from 5-(4-fluorophenyl)-3-methyl-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone and (S)-1-phenyl-1,3-propanediamine according to the General 30 Procedure. The reaction was done at 190°C for 2.5 h. MS (m/z): 430.2 (M+H)⁺; C₂₅H₂₄FN₅O requir. 429.5 (free base).

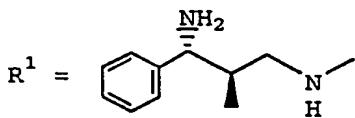


2-37 2-(((R)-3-amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride was prepared from 5-(4-fluorophenyl)-3-

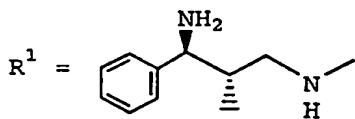
5 methyl-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone and (R)-1-phenyl-1,3-propanediamine according to the General Procedure. The reaction was done at 190°C for 3.5 h. MS (*m/z*) : 430.7 ($M+H$)⁺; $C_{25}H_{24}FN_3O$ requir. 429.5 (free base).



10 2-38 2-(((2R,3R)-3-Amino-2-methyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride was prepared from 5-(4-fluorophenyl)-3-methyl-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone and (2R,3R)-2-methyl-3-phenyl-1,3-propanediamine according to the General Procedure. The reaction was done at 190°C for 3 h. MS (*m/z*) : 444.5 ($M+H$)⁺; $C_{26}H_{26}FN_3O$ requir. 443.5 (free base).

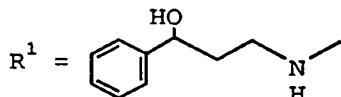


20 2-39 2-(((2S,3S)-3-Amino-2-methyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride was prepared from 5-(4-fluorophenyl)-3-methyl-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone and (2S,3S)-2-methyl-3-phenyl-1,3-propanediamine according to the General Procedure. The reaction was done at 190°C for 2 h. MS (*m/z*) : 444.4 ($M+H$)⁺; $C_{26}H_{26}FN_3O$ requir. 443.5 (free base).



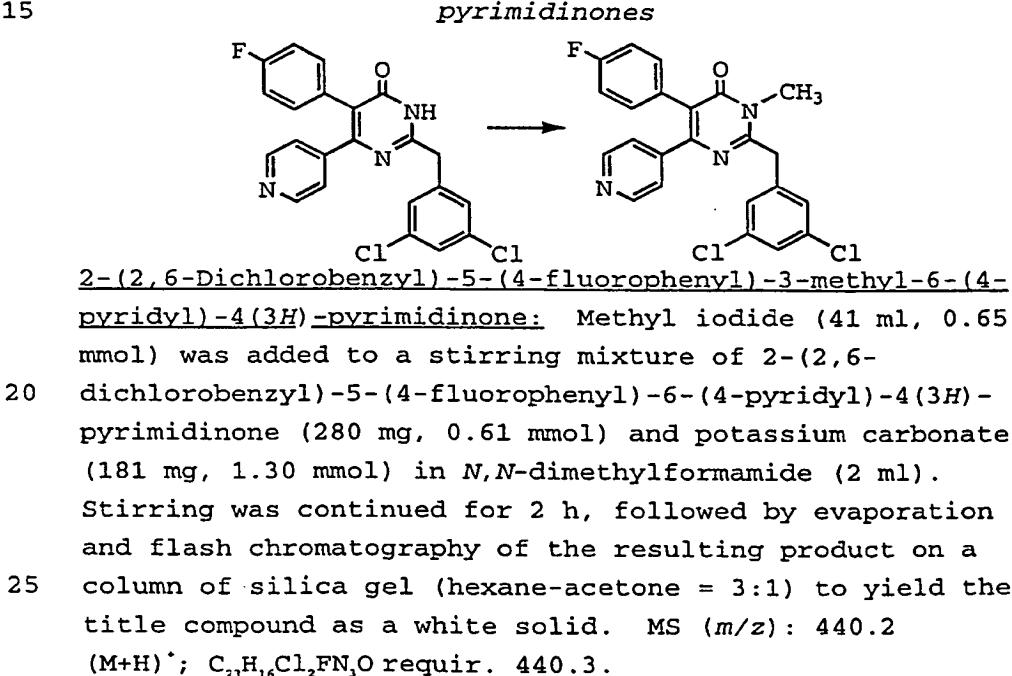
Analogously, the isomers 2-(((2S,3R)-3-Amino-2-methyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone and 2-(((2R,3S)-3-amino-2-methyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone may be prepared from the corresponding diamines.

2-40 5-(4-Fluorophenyl)-2-((-3-hydroxy-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: The reaction was done at 190°C for 3 h. MS (*m/z*): 431.2 (M+H)⁺; C₂₅H₂₃FN₄O₂ requir. 430.5.



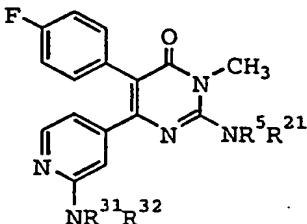
Example 3

Procedure for the preparation of N-substituted pyrimidinones



Example 4

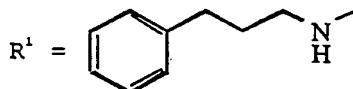
General procedure for the preparation of 2-N and 2'-N substituted 2-amino-5-(4-fluorophenyl)-3-methyl-6-(4-(2-amino)pyridyl))-4(3H)-pyrimidinones



5

Step A. 5-(4-Fluorophenyl)-3-methyl-2-methylthio-6-(4-(2-acetamido)pyridyl))-4(3H)-pyrimidinone: To a solution of 5-(4-fluorophenyl)-6-(4-(2-acetamido)pyridyl)-2-thiouracil (600 mg, 1.68 mmol) in DMF (35 mL) was added powdered sodium hydride (60% oil dispersion, 221 mg, 5.56 mmol) over 1 minute at 23°C. After 45 min, iodomethane (210 ml, 3.37 mmol) was added dropwise. After 45 min, the reaction was concentrated in vacuo (rotovap connected to high vac with a bath temperature no greater than 40°C). The residue was applied immediately to flash chromatography purification (step gradient hexane:acetone 4:1; then 3:1; then 2:1; then 1:1) to afford the desired product.

20 **Step B.** 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-amino)pyridyl))-4(3H)-pyrimidinone: A neat mixture of 5-(4-Fluorophenyl)-3-methyl-2-methylthio-6-(4-(2-acetamido)pyridyl))-4(3H)-pyrimidinone (50 mg, 0.13 mmol) and 3-phenyl-1-propylamine (88 mg, 0.65 mmol) was warmed to 190°C for 17 h. After cooling to 23°C, the reaction mixture was applied directly to purification via flash chromatography (step gradient 1%MeOH:CHCl₃ then 2%, then 3%; then 4%; then 5%) to afford the desired product: MS 30 (m/z) 430 (M+H)⁺.



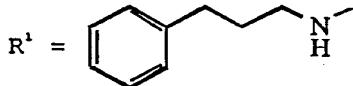
$R^{31} = \text{H}$

$R^{32} = \text{H}$

The following compounds were prepared using the
5 above procedure outlined above and an appropriate amine:

4-1 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-
amino)-6-(4-(2-acetamido)pyridyl))-4(3H)-pyrimidinone:

To a solution of 5-(4-Fluorophenyl)-3-methyl-2-((3-
phenylpropyl)-amino)-6-(4-(2-amino)pyridyl))-4(3H)-
10 pyrimidinone (11 mg, 0.026 mmol) in 600 μl of pyridine
was added (5 μl , 0.064 mmol) of acetyl chloride at 23 C.
After 2 h, the reaction was quenched with water (5 μl)
and the reaction was concentrated under a stream of
nitrogen. The reaction mixture was applied directly to
15 purification via flash chromatography (step gradient
1%MeOH:CHCl₃ then 2%, then 3%) to afford the title
compound: MS (m/z) 472 (M+H)⁺.

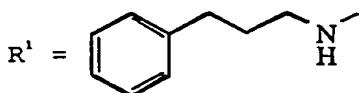


$R^{32} = \text{H}$

20 $R^{31} = \text{Ac}$

4-2 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-
amino)-6-(4-(2-methoxyacetamido)pyridyl))-4(3H)-
pyrimidinone: To a solution of 5-(4-Fluorophenyl)-3-

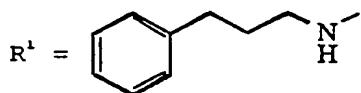
methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-
25 amino)pyridyl))-4(3H)-pyrimidinone (11 mg, 0.026 mmol)
in 600 μl of pyridine was added (5 μl , 0.064 mmol) of
methoxyacetyl chloride at 23 C. After 2 h, the reaction
was quenched with water (5 μl) and the reaction was
concentrated under a stream of nitrogen. The reaction
30 mixture was applied directly to purification via flash
chromatography (step gradient 1%MeOH:CHCl₃ then 2%, then
3%) to afford the title compound: MS (m/z) 502 (M+H)⁺.



$R^{32} = \text{H}$

$R^{31} = \text{C}(\text{O})\text{CH}_2\text{OMe}$

5 4-3 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-acetoxyacetamido)pyridyl))-4(3H)-pyrimidinone: The reaction was done in the manner of the above substituting acetoxyacetyl chloride for acetyl chloride to afford the title compound after chromatography: MS (m/z) 530 (M+H)+.



10

$R^{32} = \text{H}$

$R^{31} = \text{C}(\text{O})\text{CH}_2\text{OAc}$

15 4-4 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-hydroxyacetamido)pyridyl))-4(3H)-pyrimidinone: To a solution of 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-acetoxyacetamido)pyridyl))-4(3H)-pyrimidinone (2 mg, 0.003 mmol) in 900 μl methanol: 100 μl water was added potassium carbonate (4 mg, 0.032 mmol) as a solid at 23 C. After 3 h, the reaction was concentrated under a stream of nitrogen. The reaction mixture was diluted with chloroform (20 mL), dried (Na_2SO_4), and concentrated to afford the title compound: MS (m/z) 488 (M+H)+.

20

25 4-5 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-methylsulfonamido)pyridyl))-4(3H)-pyrimidinone: To a solution of 5-(4-Fluorophenyl)-3-

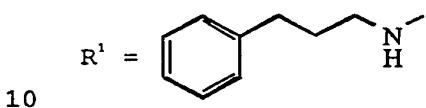
25

$R^{32} = \text{H}$

$R^{31} = \text{C}(\text{O})\text{CH}_2\text{OH}$

30 4-5 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-methylsulfonamido)pyridyl))-4(3H)-pyrimidinone: To a solution of 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-

amino)pyridyl))-4(3H)-pyrimidinone (11 mg, 0.026 mmol) in 600 μ l of pyridine was added methanesulfonyl chloride (4 μ l, 0.051 mmol) at 23 C. After 2 h, the reaction was quenched with water (5 μ l) and the reaction 5 was concentrated under a stream of nitrogen. The reaction mixture was applied directly to purification via flash chromatography (step gradient 1%MeOH:CHCl₃ then 2%) to afford the title compound: MS (m/z) 508 (M+H)⁺.

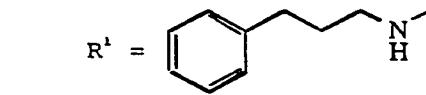


$R^{32} = \text{H}$
 $R^{31} = \text{SO}_2\text{Me}$

4-6 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-benzylamino)pyridyl))-4(3H)-pyrimidinone:

15 To a solution of 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-amino)pyridyl))-4(3H)-pyrimidinone (11 mg, 0.026 mmol) in 600 μ l of 1,2-dichloroethane was added benzaldehyde (8.9 mg, 0.084 mmol) and sodium triacetoxyborohydride (14.8 mg, 0.070 mmol) at 23 C. After 16 h, the reaction was quenched with water (15 μ l) and the reaction was concentrated under a stream of nitrogen. The reaction mixture was applied directly to purification via flash chromatography (step gradient 1%MeOH:CHCl₃ then 2%, then 3%; then 4%; then 5%) to afford the title compound: 20 MS (m/z) 458 (M+H)⁺.

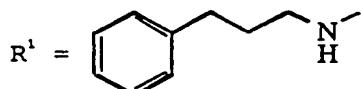
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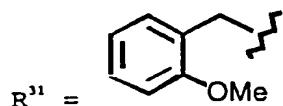
$R^{32} = \text{H}$
 $R^{31} = \text{CH}_2\text{Ph}$

30 4-7 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-(2-methoxyphenyl)methylamino)pyridyl))-4(3H)-pyrimidinone: The reaction was done in the manner

of the above substituting 2-methoxybenzaldehyde for benzaldehyde to afford the title compound after chromatography: MS (*m/z*) 550 (M+H)⁺.

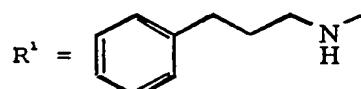


5 $R^{32} = H$



4-8 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-ethylamino)pyridyl))-4(3*H*)-pyrimidinone:

The reaction was done in the manner of the above
 10 substituting acetaldehyde for benzaldehyde to afford the title compound after chromatography: MS (*m/z*): 458 (M+H)⁺.

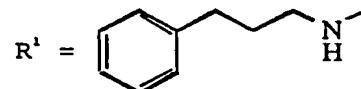


$R^{32} = H$

15 $R^{31} = Et$

4-9 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-(di-(3-methylbutyl)amino)pyridyl))-4(3*H*)-pyrimidinone: The reaction was done in the manner of the above substituting isovaleradehyde for benzaldehyde

20 to afford the title compound after chromatography: MS (*m/z*): 570 (M+H)⁺.

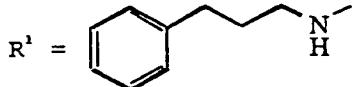


$R^{32} = CH_2CH_2CH(CH_3)_2$

$R^{31} = CH_2CH_2CH(CH_3)_2$

25 4-10 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-diethylamino)pyridyl))-4(3*H*)-pyrimidinone: The reaction was done in the manner of the above substituting acetaldehyde for benzaldehyde to

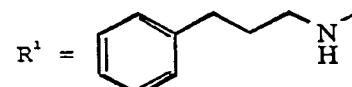
afford the title compound after chromatography: MS
 (m/z) : 486 ($M+H$)⁺.



$R^{32} =$ Et

5 $R^{31} =$ Et

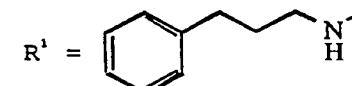
4-11 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-phenylaminocarbonyl-amino)pyridyl))-4(3H)-pyrimidinone: To a solution of 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-amino)pyridyl))-4(3H)-pyrimidinone (11 mg, 0.026 mmol) in 600 μ l of dioxane was added phenyl isocyanate (3.3 mg, 0.03 mmol) at 23°C. After 16 h, the reaction was quenched with water (15 μ l) and the reaction was concentrated under a stream of nitrogen. The reaction mixture was applied directly to purification via flash chromatography (step gradient 1%MeOH:CHCl₃ then 2%, then 3%; then 4%; then 5%) to afford the title compound: MS
 (m/z) 549 ($M+H$)⁺.



20 $R^{32} =$ H

$R^{31} =$ NH(CO)NHPH

4-12 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-methylaminocarbonyl-amino)pyridyl))-4(3H)-pyrimidinone: The reaction was done in the manner of the above substituting methylisocyanate for phenylisocyanate to afford the title compound after chromatography: MS (m/z) : 487 ($M+H$)⁺.

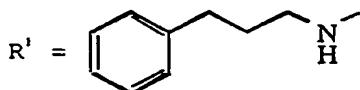


$R^{32} =$ H

30 $R^{31} =$ NH(CO)NHMe

4-13 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-(2'-amino-1'-oxo-ethylamino)pyridyl))-4(3H)-pyrimidinone:

General Procedure for mixed anhydride coupling - Isobutyl chloroformate (32 ml, 0.24 mmol) was added dropwise to a -20-30°C solution of N- α -t-Boc-glycine (5.6 mg, 0.05 mmol) and pyridine (0.6 mL). After 20 min at -20-30°C, 5-(4-fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-amino)pyridyl))-4(3H)-pyrimidinone (11 mg, 0.026 mmol) and pyridine (0.6 mL) was added in one portion. The reaction was allowed to warm to 23°C. After 16 h at 23°C, the reaction was poured into saturated bicarbonate (20 mL), extracted with ethyl acetate (2 x 50 mL), washed with brine (1 x 50 mL), and dried (Na_2SO_4). The reaction mixture was applied to purification via flash chromatography (step gradient 1%MeOH:CHCl₃ then 2%, then 3%; then 4%; then 5%) to afford the N-Boc protected title compound. The crude title compound was obtained after treatment with 50% trifluoroacetic acid:chloroform (1 mL) for 16 h. After concentration with a stream of nitrogen, the reaction mixture was applied to purification via flash chromatography (step gradient 1%MeOH:CHCl₃ then 2%, then 3%; then 4%; then 5%) to afford the title compound: MS (m/z): 487 (M+H)⁺.

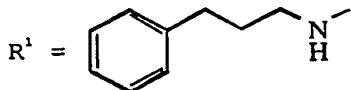


$R^{32} = \text{H}$

$R^{31} = \text{NH}(\text{CO})\text{CH}_2\text{NH}_2$

4-14 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-(4'-amino-1'-oxo-butylamino)pyridyl))-4(3H)-pyrimidinone:

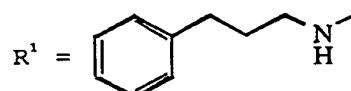
The reaction was done in the manner of the above with the following substitution: N- α -t-Boc-g-aminobutyric acid was used in place of N- α -t-Boc-glycine which after deprotection as above afforded the title compound: MS (m/z): 515 (M+H)⁺.



$R^{32} = H$

$R^{31} = NH(CO)CH_2CH_2CH_2NH_2$

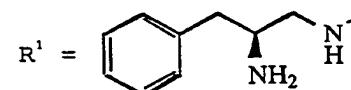
5 4-15 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-(3'-amino-1'-oxo-propylamino)pyridyl))-4(3H)-pyrimidinone: The reaction was done in the manner of the above with the following substitution: N-t-Boc- β -alanine was used in place of N- α -t-Boc-glycine which after deprotection as above afforded the title compound:
10 MS (*m/z*): 501 (M+H)⁺.



$R^{32} = H$

$R^{31} = NH(CO)CH_2CH_2NH_2$

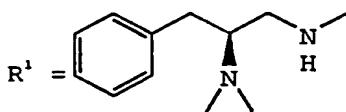
15 4-16 2-(((S)-2-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-(2-aminopyridyl))-4(3H)-pyrimidinone hydrochloride: The reaction was done at 190°C for 6 h in the above manner with the following substitution of (S)-1, 2-diamino-3-phenylpropane for 3-phenyl-1-propylamine: MS (*m/z*): 445 (M+H)⁺;



20 $R^{31} = H$

$R^{32} = H$

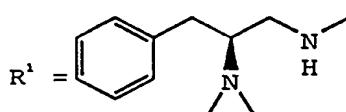
4-17 2-(((S)-2-Dimethylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-(2-aminopyridyl))-4(3H)-pyrimidinone hydrochloride: The reaction was done at 190°C for 6 h in the above manner with the following substitution of 1-amino- 2(S)-dimethylamino-3-phenylpropane for 3-phenyl-1-propylamine: MS (*m/z*): 473 (M+H)⁺;



$R^{22} = \text{H}$

$R^{21} = \text{H}$

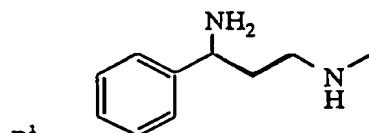
4-18 2-(((S)-2-Dimethylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-(2-acetamidopyridyl))-4(3H)-pyrimidinone hydrochloride: The reaction was done in the manner of example XX substituting 2-(((S)-2-Dimethylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-(2-aminopyridyl))-4(3H)-pyrimidinone hydrochloride for 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-amino)pyridyl))-4(3H)-pyrimidinone which afforded the title compound: MS (*m/z*): 515 (M+H)⁺;



15 $R^{22} = \text{H}$

$R^{21} = \text{Ac}$

4-19 2-(((R,S)-3-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-(2-aminopyridyl))-4(3H)-pyrimidinone hydrochloride: The reaction was done at 190°C for 12 h in the above manner with the following substitution of (3 R,S)-1,3-diamino-3-phenylpropane for 3-phenyl-1-propylamine: MS (*m/z*): 445 (M+H)⁺;



25 $R^{22} = \text{H}$

$R^{21} = \text{H}$

4-20 5-(4-Fluorophenyl)-3-methyl-2-(phenylmethylamino)-6-(4-(2-(3'-phenyl-1'-oxo-propylamino)pyridyl))-4(3H)-pyrimidinone: A neat mixture of 5-(4-fluorophenyl)-3-methyl-2-methylthio-6-(4-(2-acetamido)pyridyl))-4(3H)-pyrimidinone (260 mg, 0.13

120

mmol) and benzylamine (88 mg, 2.71 mmol) was warmed to 190 C for 17 h. After cooling to 23 C, the reaction mixture was applied directly to purification via flash chromatography (step gradient 1%MeOH:CHCl₃ then 2%, then 5% then 4%; then 5%) to afford 5-(4-Fluorophenyl)-3-methyl-2-(phenylmethylamino)-6-(4-(2-amino)pyridyl))-4(3H)-pyrimidinone. The 5-(4-fluorophenyl)-3-methyl-2-(phenylmethylamino)-6-(4-(2-amino)pyridyl))-4(3H)-pyrimidinone was converted in the manner of the above substituting hydrocinnamoyl chloride for acetyl chloride and 5-(4-fluorophenyl)-3-methyl-2-(phenylmethylamino)-6-(4-(2-amino)pyridyl))-4(3H)-pyrimidinone for 5-(4-fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-(2-amino)pyridyl))-4(3H)-pyrimidinone to afford the title compound after chromatography: MS (m/z) 534 (M+H)⁺.

R¹ = NHCH₂Ph

R³² = H

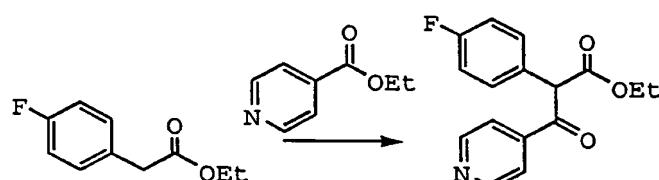
R³¹ = (CO)CH₂CH₂Ph

20

Example 5

*General procedure for the preparation of
5-(4-fluorophenyl)-6-(4-pyridyl)-2-thioalkyl-4(3H)-
pyrimidinones*

25 Step A. Ethyl 2-(4-fluorophenyl)-3-oxo-3-(4-pyridyl)-
propionate:



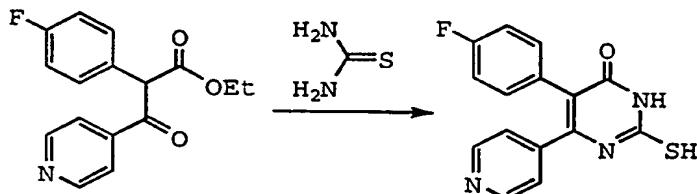
(According to: Legrand and Lozac'h, *Bull. Soc. Chim. Fr.*, 79-81 (1955)).

30 A mixture of ethyl 4-fluorophenylacetate (13 g, 71.35 mmol), ethyl isonicotinate (10.7 ml, 71.4 mmol) and sodium spheres (1.64 g, 71.34 mmol) was heated at 90-95°C under argon. The mixture started to reflux and

gradually turned into a solid. After 2.5 h, the mixture was neutralized with dil. acetic acid with cooling followed by extraction with dichloromethane. The organic solution was washed with water, dried and evaporated. Flash chromatography on a column of silica gel (hexane-acetone = 4:1, 3:1, 2:1) provided the title compound as an oil. MS (*m/z*): 287.8 (M+H)⁺; C₁₆H₁₄FNO, requir. 287.3 ¹H-NMR (CDCl₃), (ketone : enole = 1 : 0.33): δ 13.50 (s, 0.3H, OH-E), 8.81 (m, 2H, Pyrid.-K), 10 8.48 (m, 0.66 H, Pyrid.-E), 7.72 (m, 2H, Pyrid.-K), 7.38 (m, 2H, PhF-K), 7.14-7.04 (m, 2H, PhF-K; -0.65H, Pyrid.-E; -0.65H, PhF-E), 6.96 (t, 0.64H, PhF-E), 5.51 (s, 1H, CH-K), 4.23-4.2- (m, CH₂-K,E), 1.26 (t, CH₃-K,E).

Step B. 5-(4-fluorophenyl)-6-(4-pyridyl)-2-thiouracil:

15



A stirred mixture of ethyl 2-(4-fluorophenyl)-3-oxo-3-(4-pyridyl)-propionate (22.3 g, 77.6 mmol) and thiourea (5.9 g, 77.6 mmol) was reacted at 190°C under argon for 40 min. The reaction mixture was allowed to reach room temperature, taken up in acetone and the precipitate was filtered to provide the title compound. MS (*m/z*): 300.2 (M+H)⁺; C₁₅H₁₀FNOS requir. 299.3 ¹H-NMR (DMSO-d₆): δ 12.74, 12.65 (2s, 2H), 8.51 (m, 2H, Pyrid.), 7.26 (m, 2H, Pyrid.), 7.09 and 7.03 (2m, each 2H, PhF).

Alternatively, ethyl 2-(4-fluorophenyl)-3-oxo-3-(4-pyridyl)-propionate (2.87 g, 10 mmol) and thiourea (2.28 g, 30 mmol) were suspended in anhydrous p-xylene (50 ml) with very efficient stirring. To the mixture pyridinium 30 p-toluenesulfonate (100 mg) was added and refluxed for 12-16 h using a Dean-Stark apparatus with continuous removal of water (0.2 ml). Reaction mixture was cooled

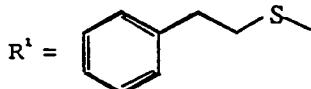
and a dark brown solid was filtered using a Buchner funnel. The collected solid was suspended in acetone (25 ml) and filtered. The acetone washed product contained a trace of thiourea, which was removed by 5 trituration with hot water (20-30 ml). The product was filtered and air dried.

Step C. General procedure:

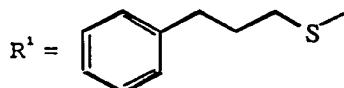
The arylalkyl bromide (0.36 mmol) was added dropwise to a stirring mixture of 5-(4-fluorophenyl)-6-(4-pyridyl)-2-thiouracil (100 mg, 0.33 mmol) and potassium carbonate (46 mg, 0.33 mmol) in *N,N*-dimethylformamide (4.6 ml). Stirring was continued for 3h followed by evaporation. Flash chromatography on a column of silica gel (hexane-acetone = 3:1, 2:1, 1:1) 15 and recrystallization from hot methanol provided the target compound.

The following compounds were obtained using the appropriate arylalkyl bromide according to the above procedure:

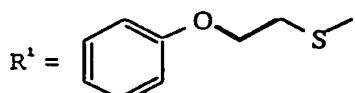
20 5-1 5-(4-Fluorophenyl)-2-(2-phenylethyl)thio-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (*m/z*): 404.2 (M+H)⁺; C₂₃H₁₈FN₃OS requir. 403.4. ¹H-NMR (DMSO-d₆): δ 13.08 (bs, 0.7H), 8.49 (m, 2H, Pyrid.), 7.30-7.06 (m, 11H, Pyrid., Ph, PhF), 3.41 (dd, 2H, CH₂S), 3.00 (t, 2H, CH₃).



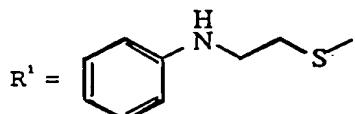
25 5-2 5-(4-Fluorophenyl)-2-(3-phenylpropyl)thio-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (*m/z*): 418.0 (M+H)⁺; C₂₄H₂₀FN₃OS requir. 417.5. ¹H-NMR (DMSO-d₆): δ 13.10 (bs, 0.7H), 8.47 (m, 2H, Pyrid.), 7.29-7.06 (m, 11H, Pyrid., Ph, PhF), 3.18 (t, 2H, CH₂S), 2.71 (t, 2H, CH₂Ph), 2.03 (m, 2H, CH₂).



5-3 5-(4-Fluorophenyl)-2-(2-phenoxyethyl)thio-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (*m/z*): 420.0 (M+H)⁺; C₂₃H₁₈FN₂O₂S requir. 419.5. ¹H-NMR (DMSO-d₆): d 13.20 (bs, 0.7H), 8.46 (m, 2H, Pyrid.), 7.24-7.07 (m, 8H, Pyrid., PhF, Ph), 6.95 (d, 2H, Ph), 6.92 (t, overlapped, 1H, Ph), 4.30 (t, 2H, CH₂O), 3.58 (t, 2H, CH₂S).



5-4 5-(4-Fluorophenyl)-2-(2-phenylaminoethyl)thio-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (*m/z*): 419.0 (M+H)⁺; C₂₃H₁₉FN₂OS requir. 418.5. ¹H-NMR (DMSO-d₆): d 13.20 (bs, 0.8H), 8.48, 7.22 (2m, each 2H, Pyrid.), 7.16, 7.10 (2m, each 2H, PhF), 6.89 (t, 2H, Ph), 6.54 (d, 2H, Ph), 6.48 (t, 1H, Ph), 5.90 (bs, 0.6H, NH), 3.43-3.25 (m, 2CH₂).

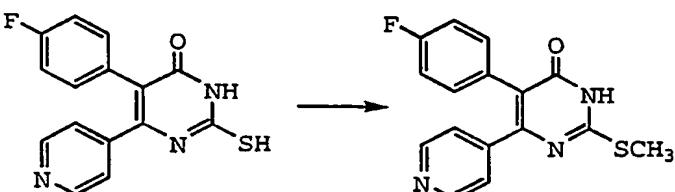


15

Example 6

General procedure for the preparation of 2-N substituted 2-amino-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinones:

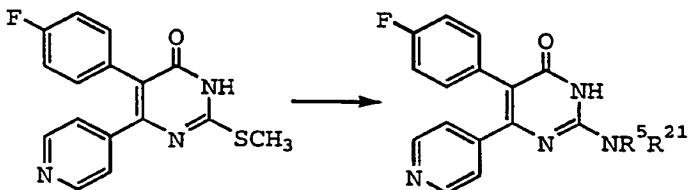
Step A. 5-(4-Fluorophenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone:



Methyl iodide (90 ml, 1.44 mmol) was added dropwise to a stirred mixture of 5-(4-fluorophenyl)-6-(4-pyridyl)-2-thiouracil (430 mg, 1.44 mmol) and potassium carbonate (198 mg, 1.43 mmol) in *N,N*-dimethylformamide (13 ml) at ice-bath temperature. After 40 min, it was evaporated and the crude product purified by flash chromatography on a column of silica gel (hexane-acetone

= 2:1, 1:1, 1:2) to provide the title compound as a solid. MS (*m/z*): 314.2 ($M+H$)⁺; C₁₆H₁₂FN₃OS requir. 313.3. ¹H-NMR (DMSO-d₆): d 13.10 (bs), 8.47, 7.22 (2m, each 2H, Pyrid.), 7.16, 7.10 (2m, each 2H, PhF), 2.56 (s, 3H, 5 CH₃).

Step B. General procedure:



A mixture of 5-(4-fluorophenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone (100 mg, 0.32 mmol) and an amine HNR⁵R²¹ (1 mmol) was heated at 180°C for 2 h. The resulting product was purified by flash chromatography on a column of silica gel (hexane-acetone or methanol-dichloromethane or dichloromethane-methanol-conc. ammonium hydroxide) to provide the target compound.

The following compounds were prepared using the general procedure outlined above and an appropriate amine:

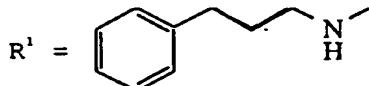
6-1 2-(2-Chlorophenyl)ethyl-amino)-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (*m/z*): 421.2 ($M+H$)⁺; C₂₃H₁₈ClFN₄O requir. 420.9. ¹H-NMR (DMSO-d₆): d 11.24 (bs), 8.44, 7.16 (2m, each 2H, Pyrid.), 7.43, 7.38 (2dd, each 1H, PhCl), 7.30, 7.26 (2dt, each 1H, PhCl), 7.10-7.00 (m, 2H, PhF), 6.74 (bs, 1H, NH), 3.60 (q, 2H, CH₂N), 3.03 (t, 2H, CH₂).

R¹ =

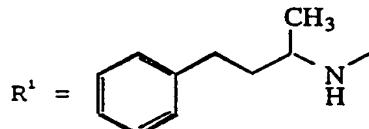
6-2 5-(4-Fluorophenyl)-2-((3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (*m/z*): 401.2 ($M+H$)⁺; C₂₄H₂₁FN₄O requir. 400.5. ¹H-NMR (DMSO-d₆): d 11.16 (bs), 8.44, 7.14 (2m, each 2H, Pyrid.), 7.32-7.01 (m, 9H, Ph,

125

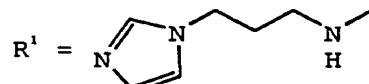
PhF), 6.78 (bs, NH), 3.36 (q, 2H, CH₂N), 2.67 (t, 2H, CH₂Ph), 1.89 (m, 2H, CH₂).



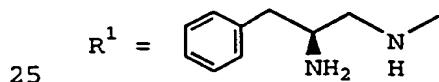
6-3 5-(4-Fluorophenyl)-2-((1-methyl-3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone: A reaction time of 15 h at 180°C was required. MS (m/z): 415.0 (M+H)⁺; C₂₃H₂₃FN₄O requir. 414.5. ¹H-NMR (CDCl₃): d 8.48 (m, 2H, Pyrid.), 7.28-7.08 (m, 9H, Pyrid., Ph, PhF), 6.94 (m, 2H, PhF), 5.67 (bs, 1H, NH), 4.08 (m, 1H, CHCH₃), 2.61 (t, 2H, CH₂Ph), 1.67 (m, 2H, CH₂), 1.08 (d, 3H, CH₃).



6-4 5-(4-Fluorophenyl)-2-((3-imidazolylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (m/z): 391.0 (M+H)⁺; C₂₁H₁₉FN₆O requir. 390.4. ¹H-NMR (DMSO-d₆): d 11.24 (bs), 8.42, 7.12 (2m, each 2H, Pyrid.), 7.62, 7.18 (2s, each 1H, Imid.), 7.08-6.99 (m, 4H, PhF), 6.88 (s, 1H, Imid.), 4.02 (t, 2H, CH₂N), 3.28 (overlapped by water signal, CH₂NH), 2.00 (m, 2H, CH₂).

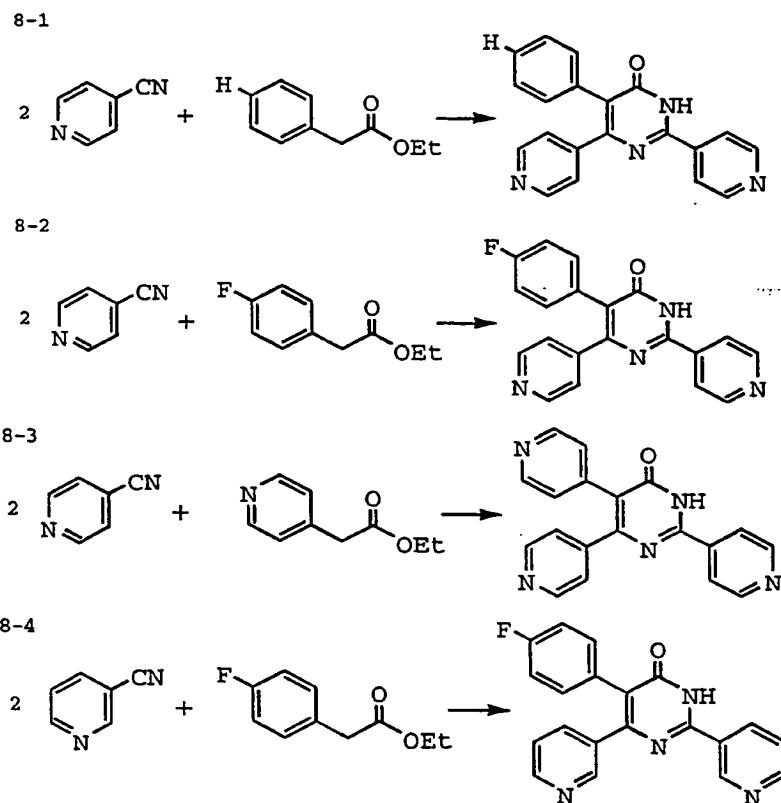


6-5 2-(((S)-2-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: The reaction was done at 170°C for 7 h. MS (m/z): 416.1 (M+H)⁺; C₂₆H₂₂FN₅O requir. 415.5.



Example 7*5-(4-Fluorophenyl)-2-hydrazino-6-(4-pyridyl)-4(3H)-pyrimidinone*

A mixture of 5-(4-fluorophenyl)-6-(4-pyridyl)-2-thiouracil (500 mg, 1.66 mmol) and hydrazine hydrate (800 ml, ~14 mmol) was heated at 120°C for 60 min. It was evaporated and the reaction product was recrystallized from hot methanol to provide the title compound. MS (*m/z*): 298.0 (M+H)⁺; C₁₅H₁₂FN₄O requir.
 10 297.3. ¹H-NMR (DMSO-d₆): δ 8.41, 7.12 (2m, each 2H, Pyrid.), 7.05, 7.00 (2m, each 2H, PhF).
 R¹ = NH-NH₂

Example 8*General procedure for the preparation of 5-aryl-2,6-dipyridyl-(3H)-pyrimidinones*

These compounds were prepared according to the literature (Kabbe, *supra*; German Patent 1271116 (1968)) as follows:

A stirred mixture of the ethyl phenylacetate (3.13 mmol), cyanopyridine (6.24 mmol) and sodium methoxide (3.5 mmol) in n-butanol (1.2 ml) was heated at 110°C for 2h. The reaction mixture was concentrated and dissolved in water (4 ml), followed by the addition of aqueous sat. ammonium chloride (2 ml). The precipitate was 10 filtered and recrystallized from hot methanol.

The following compounds were prepared according to this procedure using the appropriate starting materials:

8-1 5-Phenyl-2,6-bis-(4-pyridyl)-4-(3H)-pyrimidinone: MS (m/z): 327.2 (M+H)⁺; C₂₀H₁₄N₄O requir. 326.4. ¹H-NMR (DMSO-d₆): d 8.78, 8.47, 8.13 (3m, each 2H, Pyrid.), 7.40-7.14 (m, 7H, Ph, Pyrid.).

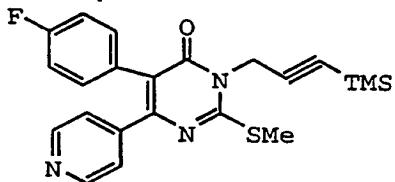
8-2 5-(4-Fluorophenyl)-2,6-bis-(4-pyridyl)-4(3H)-pyrimidinone: MS (m/z): 345.2 (M+H)⁺; C₂₀H₁₃FN₄O requir. 344.4 ¹H-NMR (DMSO-d₆): d 8.80, 8.49, 8.13 (3m, each 2H, Pyrid.), 7.40-7.08 (m, 6H, PhF, Pyrid.).

8-3 2,5,6-Tris-(4-pyridyl)-4(3H)-pyrimidinone was prepared according to the general procedure by reacting ethyl 4-pyridylacetate and 4-cyanopyridine in the presence of sodium methoxide. MS (m/z): 328.2 (M+H)⁺; C₁₉H₁₃N₃O requir. 327.4 ¹H-NMR (DMSO-d₆): 8.65, 8.45, 8.35, 8.18, 7.25, 7.13 (6m, each 2H, Pyrid.).

8-4 5-(4-Fluorophenyl)-2,6-bis-(3-pyridyl)-4(3H)-pyrimidinone: MS (m/z): 345.2 (M+H)⁺; C₂₀H₁₃FN₄O requir. 344.4 ¹H-NMR (DMSO-d₆): d 9.34, 8.77, 8.54, 8.48, 7.78, 7.60, 7.34 (7m, 3x1H, 2H, 3x1H, Pyrid.), 7.26, 7.15 (2m, each 2H, PhF).

Example 9

3-(3-trimethylsilyl-2-propynyl)-5-(4-fluorophenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone

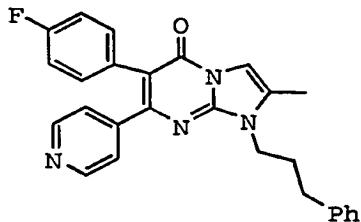


5 The preparation of the title compound was carried out in the same manner as 3-ethyl-5-(4-fluorophenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone with the following substitution: 3-bromo-1-(trimethylsilyl)-1-propyne was used in place of ethyl bromide.

10

Example 10

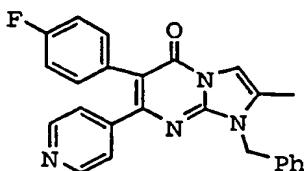
6-(4-Fluorophenyl)-2-methyl-1-(3-phenylpropyl)-7-pyridin-4-yl-1H-imidazo(1,2-a)pyrimidin-5-one



15 A neat mixture of 3-(3-trimethylsilyl-2-propynyl)-5-(4-fluorophenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone (50 mg, 0.12 mmol) and 3-phenyl-1-propylamine (67 mg, 0.47 mmol) was warmed to 190°C for 17 h. After cooling to 23°C, the reaction mixture was
20 applied directly to purification via flash chromatography (step gradient 1%MeOH:CHCl₃ then 2%, then 3%;) to afford the desired product: MS (m/z) 439 (M+H)⁺.

Example 11

6-(4-Fluorophenyl)-2-methyl-1-benzyl-7-pyridin-4-yl-1H-imidazo(1,2-a)pyrimidin-5-one

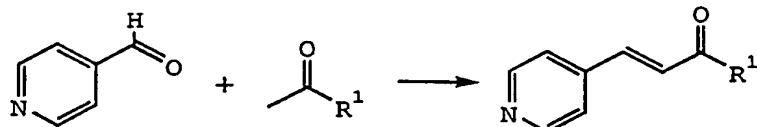


5 The preparation of the title compound was carried out in the same manner as 6-(4-Fluorophenyl)-2-methyl-1-(3-phenyl propyl)-7-pyridin-4-yl-1H-imidazo(1,2-a)pyrimidin-5-one with the following substitution:
benzylamine for 3-phenyl-1-propylamine; MS (*m/z*): 411
10 (M+H)⁺.

Example 12

General procedure for the preparation of 6-substituted 3-phenyl-4-(4-pyridyl)-2(1H)-pyridones

Step A. General procedure for the preparation of 1-aryl-3-(4-pyridyl)-2-propene-1-one :



At ice-bath temperature, piperidine (206 ml), acetic acid (206 ml) and 4-pyridinecarboxaldehyde (1.6 ml, 16.6 mmol) were mixed. Then the acetophenone (12.0 mmol) was added at room temperature and the mixture was heated at 130°C for 1.5 h. The reaction mixture was diluted with dichloromethane, washed with aqueous sodium hydrogen carbonate and water followed by drying and evaporation. The crude product was purified by column chromatography on silica gel (hexane-acetone = 3:1).

The following compounds were prepared according to this procedure using the appropriate acetophenone derivative:

1-Phenyl-3-(4-pyridyl)-2-propene-1-one: MS (*m/z*): 210.1
30 (M+H)⁺; C₁₄H₁₁NO requir. 209.3.

130

1-(4-Methylphenyl)-3-(4-pyridyl)-2-propene-1-one: MS
 (m/z) : 224.2 ($M+H$)⁺; $C_{15}H_{13}NO$ requir. 223.3.

1-(4-Ethylphenyl)-3-(4-pyridyl)-2-propene-1-one: MS
 (m/z) : 237.8 ($M+H$)⁺; $C_{16}H_{15}NO$ requir. 237.3.

5 1-(4-Isopropylphenyl)-3-(4-pyridyl)-2-propene-1-one: MS
 (m/z) : 252.0 ($M+H$)⁺; $C_{17}H_{15}NO$ requir. 251.3.

1-(2-Methylphenyl)-3-(4-pyridyl)-2-propene-1-one: MS
 (m/z) : 223.8 ($M+H$)⁺; $C_{15}H_{13}NO$ requir. 223.3.

10 1-(2,4-Dimethylphenyl)-3-(4-pyridyl)-2-propene-1-one: MS
 (m/z) : 238.0 ($M+H$)⁺; $C_{16}H_{15}NO$ requir. 237.3.

1-(2-Methoxyphenyl)-3-(4-pyridyl)-2-propene-1-one: MS
 (m/z) : 240.0 ($M+H$)⁺; $C_{15}H_{13}NO_2$ requir. 239.3

1-(4-Chlorophenyl)-3-(4-pyridyl)-2-propene-1-one: MS
 (m/z) : 244.0 ($M+H$)⁺; $C_{14}H_{10}ClNO$ requir. 243.7.

15 1-(4-Cyanophenyl)-3-(4-pyridyl)-2-propene-1-one: MS
 (m/z) : 235.1 ($M+H$)⁺; $C_{15}H_{10}N_2O$ requir. 234.3.

1-(a-Naphthyl)-3-(4-pyridyl)-2-propene-1-one: MS (m/z) :
260.0 ($M+H$)⁺; $C_{18}H_{13}NO$ requir. 259.3.

1,3-Bis-(4-pyridyl)-2-propene-1-one: MS (m/z) : 211.0
20 ($M+H$)⁺; $C_{13}H_{10}N_2O$ requir. 210.2.

3-(4-Pyridyl-1-(2-thienyl)-2-propene-1-one: MS (m/z) :
216.0 ($M+H$)⁺; $C_{12}H_9NOS$ requir. 215.3.

1-(2-Furyl-3-(4-pyridyl)-2-propene-1-one: MS (m/z) :
200.0 ($M+H$)⁺; $C_{12}H_9NO_2$ requir. 199.2.

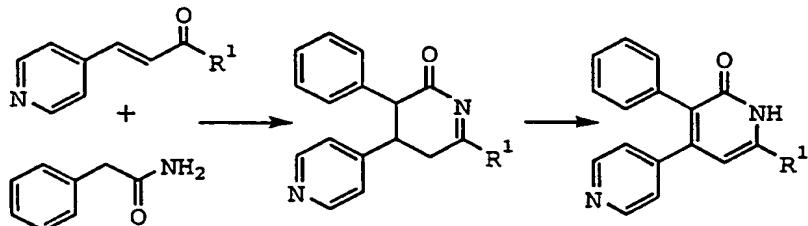
25 1-Cyclohexyl-3-(4-pyridyl)-2-propene-1-one was prepared
in the same way using acetyl cyclohexane: MS (m/z) :
216.2 ($M+H$)⁺; $C_{14}H_{11}NO$ requir. 215.3.

1-tert-Butyl-3-(4-pyridyl)-2-propene-1-one: A mixture of
3,3-dimethyl-2-butanone (2.5 ml, 20.0 mmol), 4-

30 pyridinecarboxaldehyde (2.15 ml, 22.3 mmol), ethanol
(7.6 ml), and 4.5% aqueous sodium hydroxide (4.6 ml) was
kept at room temperature for 12 h. It was diluted with
dichloromethane, washed with aqueous hydrochloric acid
and water, dried and evaporated. Subsequent column

35 chromatography (hexane - ethyl acetate = 3:1) provided
the title compound. MS (m/z) : 190.4 ($M+H$)⁺; $C_{12}H_{13}NO$
requir. 189.3.

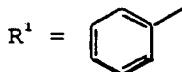
Step B. General procedure for the preparation of 6-substituted 3-phenyl-4-(4-pyridyl)-2(1H)-pyridones:



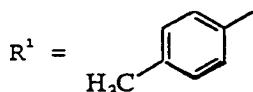
Sodium (40 mg, 1.74 mmol) was dissolved in a
 5 stirring mixture of phenylacetamide (880 mg, 6.51 mmol)
 and ethanol (5ml). If solubility allowed, the 1-
 substituted 3-(4-pyridyl)-2-propene-1-one (5.4 mmol) was
 added portionwise as an ethanolic solution (20 ml) to
 the refluxing phenylacetamide solution or it was added
 10 at room temperature as a solid. The mixture was kept
 under reflux for 1.5 h and was then allowed to reach
 room temperature. 2N Hydrochloric acid was added to a
 pH value of 5 followed by the addition of a few ml of
 15 water. The product that crystallized from this mixture
 was filtered, washed subsequently with ethanol, water,
 ethanol and recrystallized from methanol. If the
 product did not crystallize from the reaction mixture on
 addition of hydrochloric acid, then the mixture was
 20 evaporated and the remainder taken up in
 dichloromethane. The organic solution was washed with
 water, dried and evaporated. The resultant product was
 crystallized from hot acetone and recrystallized from
 methanol.

The following compounds were prepared according to
 25 this procedure using the 2-(4-pyridyl)-2-propene-1-one
 derivatives described in Example 12.a:

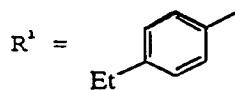
12-1 3,6-Diphenyl-4-(4-pyridyl)-2(1H)-pyridone: MS
 (m/z): 325.4 (M+H)⁺; C₂₂H₁₆N₂O requir. 324.4. ¹H-NMR (DMSO-d₆): δ 8.63 (m, 2H, Pyrid.), 7.86 (m, 2H), 7.58-7.45,
 30 7.29-7.08 (2m).



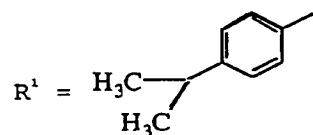
12-2 6-(4-Methylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (*m/z*): 339.2 ($M+H$)⁺; $C_{23}H_{18}N_2O$ requir. 338.4.
¹H-NMR (DMSO-d₆): d 8.44 (m, 2H, Pyrid.), 7.79 (d, 2H),
5 7.32 (d, 2H), 7.26-7.01 (m, 7H, Ph, Pyrid.), 6.67 (bs, 1H).



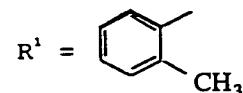
12-3 6-(4-Ethylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (*m/z*): 353.0 ($M+H$)⁺; $C_{24}H_{20}N_2O$ requir. 352.4.
10 ¹H-NMR (DMSO-d₆): d 8.42 (m, 2H, Pyrid.), 7.79 (d, 2H),
7.33 (d, 2H), 7.24-7.06 (m, 7H, Ph, Pyrid.), 6.65 (bs, 1H, CH=), 2.66 (q, 2H, CH₂), 1.21 (t, 3H, CH₃).



12-4 6-(4-Isopropylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (*m/z*): 367.0 ($M+H$)⁺; $C_{25}H_{22}N_2O$ requir. 366.5.
¹H-NMR (DMSO-d₆): d 8.45 (m, 2H, Pyrid.), 7.82 (d, 2H),
7.39 (d, 2H), 7.28-7.10 (m, 7H, Ph, Pyrid.), 6.67 (bs, 1H, CH=), 2.98 (m, 1H, CH(CH₃)₂), 1.27, 1.25 (2s, each 3H, 2CH₃).

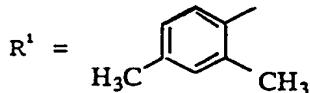


20 12-5 6-(2-Methylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (*m/z*): 339.2 ($M+H$)⁺; $C_{23}H_{18}N_2O$ requir. 338.4.
¹H-NMR (DMSO-d₆): d 8.40 (m, 2H, Pyrid.), 7.45-7.09 (m, 11H, Ph, Pyrid.), 6.21 (bs, 1H, CH=), 2.39 (s, 3H, CH₃).

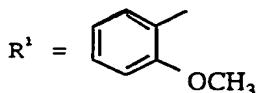


25 12-6 6-(2,4-Dimethylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (*m/z*): 353.0 ($M+H$)⁺; $C_{24}H_{20}N_2O$ requir. 352.4. ¹H-NMR (DMSO-d₆): d 8.42 (m, 2H, Pyrid.), 7.29

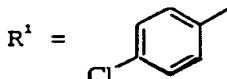
(d, 1H), 7.23-7.06 (m, 9H, Ph, Pyrid.), 6.17 (bs, 1H, CH=), 2.34, 2.31 (2s, each 3H, 2CH₃).



12-7 6-(2-Methoxyphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (m/z) : 355.0 (M+H)⁺; C₂₂H₁₈N₂O₂ requir. 354.4. ¹H-NMR (DMSO-d₆): d 8.41 (m, 2H, Pyrid.), 7.49 (bd, 1H), 7.44 (m, 1H), 7.24-7.06 (m, 8H, Ph, Pyrid.), 7.02 (dt, 1H), 6.32 (bs, 1H, CH=), 3.82 (s, 3H, CH₃).



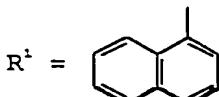
10 12-8 6-(4-Chlorophenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (m/z) : 359.2 (M+H)⁺; C₂₂H₁₅ClN₂O requir. 358.8. ¹H-NMR (DMSO-d₆): d 8.42 (m, 2H, Pyrid.), 7.93 (bd, 2H), 7.54 (m, 2H), 7.26-7.08 (m, 7H, Ph, Pyrid.), 6.80 (bs, 1H, CH=).



15 12-9 6-(4-Cyanophenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (m/z) : 350.2 (M+H)⁺; C₂₂H₁₅N₃O requir. 349.4. ¹H-NMR (DMSO-d₆): d 8.45 (m, 2H, Pyrid.), 8.16 (bd, 2H), 7.98 (d, 2H), 7.32-7.00 (m, 8H, Ph, Pyrid., CH=).



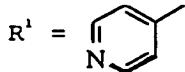
20 12-10 6-(a-Naphthyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (m/z) : 375.0 (M+H)⁺; C₂₆H₁₈N₂O requir. 374.5. ¹H-NMR (DMSO-d₆): d 8.38 (m, 2H, Pyrid.), 8.06-7.98 (m, 3H), 7.67 (dd, 1H), 7.62-7.54 (m, 3H), 7.25-7.11 (m, 7H, Ph, Pyrid.), 6.38 (bs, 1H, CH=).



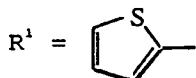
25 12-11 3-Phenyl-4,6-bis-(4-pyridyl)-2(1H)-pyridone: MS (m/z) : 326.0 (M+H)⁺; C₂₁H₁₅N₃O requir. 325.4. ¹H-NMR

134

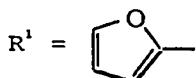
(DMSO-d₆: d 8.69, 8.43 (2m, each 2H, Pyrid.), 7.92 (bs, 2H), 7.28-7.05 (m, 8H).



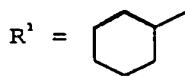
12-12 3-Phenyl-4-(4-pyridyl)-6-(2-thienyl)-2(1H)-pyridone: MS (m/z) : 331.0 (M+H)⁺; C₂₂H₁₄N₂OS requir. 330.4.
¹H-NMR (DMSO-d₆): d 8.44 (m, 2H, Pyrid.), 7.90, 7.70 (2bd, each 1H), 7.28-7.08 (m, 9H).



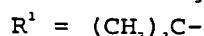
12-13 6-(2-Furyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (m/z) : 315.0 (M+H)⁺; C₂₁H₁₄N₂O, requir. 314.4.
¹H-NMR (DMSO-d₆): d 8.44 (m, 2H, Pyrid.), 7.90 (s, 1H), 7.43 (bs, 1H), 7.27-7.08 (m, 7H, Ph, Pyrid.), 6.71 (m, 2H).



12-14 6-Cyclohexyl-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (m/z) : 331.2 (M+H)⁺; C₂₂H₂₂N₂O requir. 330.4.
¹H-NMR (DMSO-d₆): d 8.40 (m, 2H, Pyrid.), 7.22-7.13, 7.10-7.03 (2m, 7H, Ph, Pyrid.), 6.04 (bs, 1H, CH=), 1.95-1.15 (m, 11H, cyclohex.).

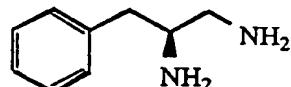


12-15 6-tert-Butyl-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone: MS (m/z) : 305.0 (M+H)⁺; C₂₀H₂₀N₂O requir. 304.4.
¹H-NMR (DMSO-d₆): d 8.39 (m, 2H, Pyrid.), 7.20-7.12, 7.10-7.02 (2m, 7H, Ph, Pyrid.), 6.02 (bs, 1H, CH=), 1.31 (s, 9H, 3CH₃).



Example 13

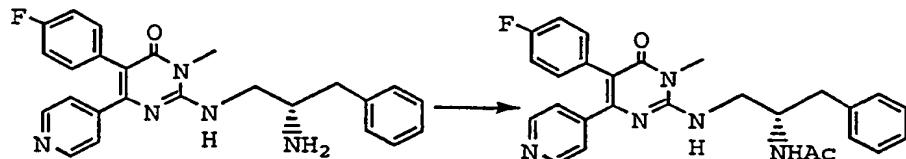
Procedure for the preparation of (S)-1,2-Benzylethylendiamine



5 **(S)-1,2-Benzylethylendiamine:** The diamine was prepared according to the literature (H. Brunner, P. Hankofer, U. Holzinger, B. Treitinger and H. Schoenenberger, Eur. J. Med. Chem. 25, 35-44, (1990)) by reduction of L-phenylalanine amide with lithium aluminium hydride. The 10 (R)-enantiomer was prepared in the same manner from D-phenylalanine amide.

Example 14

15 *Procedure for the preparation of 2-(((S)-2-Acetamido-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone*

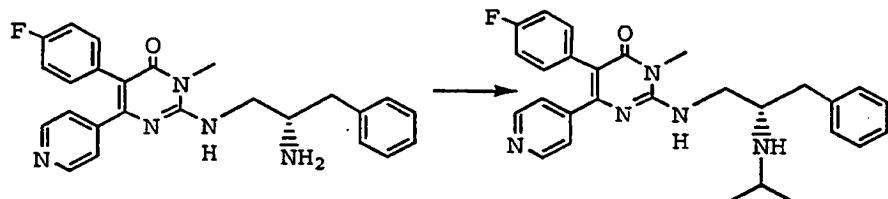
**2-(((S)-2-Acetamido-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone:**

20 A solution of 2-(((S)-2-amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone (25 mg, 0.058 mmol) and acetic anhydride (200 ml) in methanol (2 ml) was kept at room temperature for 1 h. Evaporation followed by chromatography of the resultant product on a column of silica gel (10% methanol/dichloromethane) provided the title compound.

25 MS (*m/z*): 472.3 (*M+H*)⁺; C₂₇H₂₆FN₅O₂ requir. 471.5.

Example 15

Procedure for the preparation of 5-(4-Fluorophenyl)-2-(((S*)-2-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride*

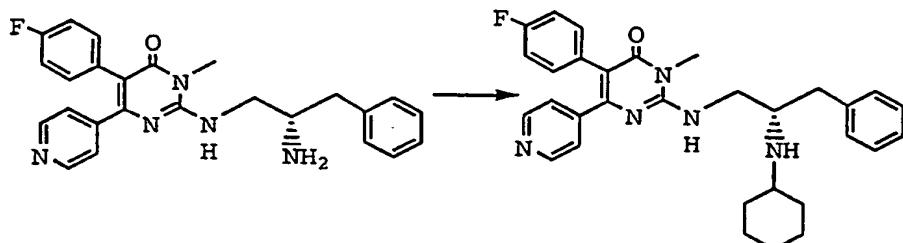


5

5-(4-Fluorophenyl)-2-((*(S*)-2-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: Sodium triacetoxymethane (23 mg, 0.109 mmol) was added to a stirring mixture of 2-((*(S*)-2-amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride (50 mg, 0.107 mmol), triethylamine (15 ml, 0.108 mmol) and acetone (7.9 ml, 0.108 mmol) in 1,2-dichloroethane (0.8 ml). After 4h, the reaction was quenched by the addition of sat. aqu. sodium hydrogencarbonate, followed by extraction with dichloromethane, drying of the organic solution and evaporation. Chromatography on a column of silica gel (10% methanol/chloroform) provided the title compound as a free base which was converted into the monohydrochloride by the addition of 4N hydrochloric acid/dioxane (21 mmol, 0.08 mmol) to its methanolic solution (1 ml) and subsequent evaporation. MS (*m/z*): 472.1 ($M+H$)⁺; C₂₈H₃₀FN₅O requir. 471.6 (free base).

Example 16

Procedure for the preparation of 5-(4-Fluorophenyl)-2-(((S)-2-N-cyclohexylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride



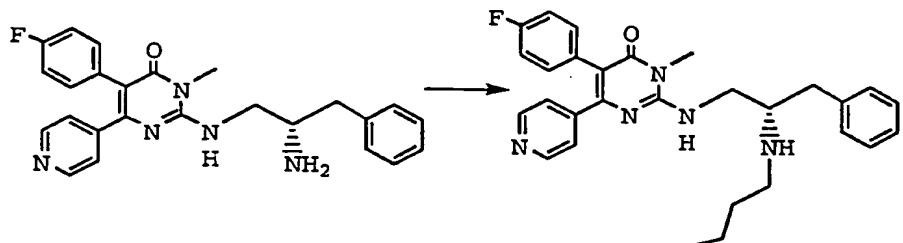
5

5-(4-Fluorophenyl)-2-(((S)-2-N-cyclohexylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: Utilizing cyclohexanone, 5-(4-fluorophenyl)-2-(((S)-2-N-cyclohexylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone was prepared in the same manner as 5-(4-fluorophenyl)-2-(((S)-2-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: MS (m/z): 511.6 (M)⁺; C₃₁H₃₄FN₅O requir. 511.6 (free base).

15

Example 17

Procedure for the preparation of 2-(((S)-2-N-n-Butylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride



20

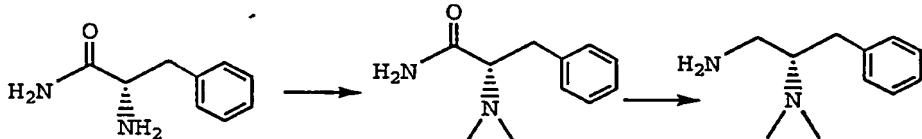
2-(((S)-2-N-n-Butylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: Sodium triacetoxyborohydride (28 mg, 0.13 mmol) was added to a stirring mixture of 2-(((S)-2-amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone (41 mg, 0.095

25

mmol) and butyraldehyde (8.5 ml, 0.094 mmol) in 1,2-dichloroethane (0.8 ml). After 2 h, the reaction was quenched by the addition of sat. aqu. sodium hydrogencarbonate, followed by extraction with dichloromethane, drying of the organic solution and evaporation. Chromatography on a column of silica gel (5% methanol/chloroform) provided the title compound as a free base which was converted into the monohydrochloride by the addition of 4N hydrochloric acid/dioxane (12 mmol, 0.048 mmol) to its methanolic solution (1 ml) and subsequent evaporation. MS (*m/z*): 486.2 (M+H)⁺; C₂₉H₃₂FN₅O requir. 485.6 (free base).

Example 18

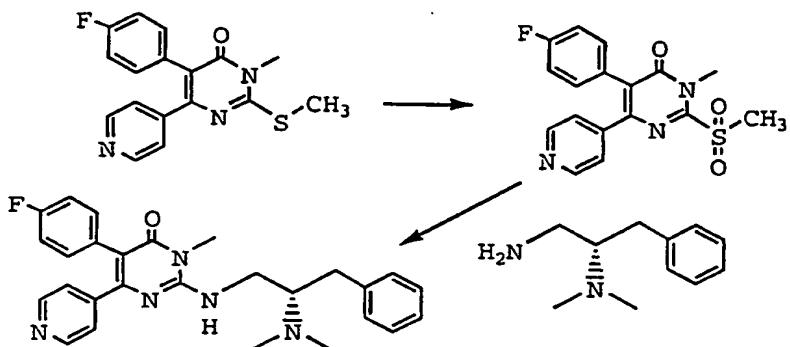
Procedure for the preparation of (S)-2-N,N-Dimethylamino-3-phenylpropylamine



(S)-2-N,N-Dimethylamino-3-phenylpropylamine: Sodium triacetoxyhydride (13.0 g, 61.3 mmol) was added to a stirring mixture of phenylalanine amide (3.6 g, 21.9 mmol) and 37% formaldehyde solution (4.4 ml, 58.7 mmol) in 1,2-dichloroethane (77 ml). After stirring for 2 h, the reaction was quenched by the addition of sat. aqu. sodium hydrogencarbonate. Then potassium hydroxide pellets were added followed by extraction with dichloromethane, drying of the organic solution and evaporation. The resulting (S)-2-N,N-dimethylamino-3-phenylpropylamide was reduced with lithium aluminium hydride according to the literature (H. Brunner, P. Hankofer, U. Holzinger, B. Treittinger and H. Schoenenberger, Eur. J. Med. Chem. 25, 35-44, (1990)) to provide the title compound.

Example 19

Procedure for the preparation of 2-(((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride



Step A. 5-(4-Fluorophenyl)-3-methyl-2-methylsulfonyl-6-(4-pyridyl)-4(3H)-pyrimidinone: A mixture of 5-(4-fluorophenyl)-3-methyl-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone (400 mg, 1.22 mmol) and Oxone® (potassium peroxyxonate, 2.3 g, 3.74 mmol) in methanol (100 ml) and water (45 ml) was stirred for 13 h. The solvent was concentrated to about 50 ml, followed by extraction with dichloromethane, drying of the organic solution and evaporation. The resulting white solid was used without purification in the next step.

Step B. 2-(((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: A mixture of crude 5-(4-fluorophenyl)-3-methyl-2-methylsulfonyl-6-(4-pyridyl)-4(3H)-pyrimidinone (430 mg g, 1.19 mmol) and (S)-2-N,N-dimethylamino-3-phenylpropylamine (600 mmol, ~3.4 mmol) was stirred at room temperature for 1h and then briefly warmed at 50°C. Column chromatography on silica gel (3-5% methanol/chloroform) provided the title compound as a free base which was converted into the monohydrochloride by the addition of 4N hydrochloric acid/dioxane (160 mmol, 0.64 mmol) to its methanolic solution (4 ml) and

subsequent evaporation. MS (*m/z*): 458.0 (M+H)⁺;
C₂₇H₂₈FN₅O requir. 457.5 (free base).

Example 20

5 *5-(4-fluorophenyl)-6-(4-(2-acetamido)-pyridyl)-2-*
thioalkyl-4(3H)-pyrimidinones

Step A. Ethyl 2-(4-fluorophenyl)-3-oxo-3-(4-(2-
acetamido)-pyridyl))-propionate:

A solution of 2-chloroisonicotinic acid (25.0g, 0.16
10 mol) in 65 mL of concentrated ammonium hydroxide was
warmed to 205 Celsius in a steel bomb for 72 h. After
cooling to 23 C, the solution was acidified to a pH of 1
using 6N HCl and subsequently filtered to remove
unreacted starting material. The solution was
15 concentrated to one fourth the original volume (approx
200 mL), in vacuo, and carefully adjusted to a pH of 6
using 1 N NaOH. After storing the cloudy solution at 0
C for 20 h, the desired 2-aminoisonicotinic acid was
filtered off. To a suspension of 2-aminoisonicotinic
20 acid in ethanol (600 mL) was added 47.1 mL of 4 N
anhydrous HCl in dioxane. After warming to achieve
reflux for 20 h, an additional 47.1 mL of 4 N anhydrous
HCl in dioxane was added and the reaction was warmed to
reflux for an additional 20 h. Concentration with a
25 stream of nitrogen in the hood was followed by further
concentration in vacuo, the remaining solid was diluted
with saturated bicarbonate (200 mL), extracted with
ethyl acetate (2 x 200mL), dried (Na₂SO₄). After
concentration in vacuo, the desired ethyl 2-
30 aminoisonicotinate was obtained. To a solution of ethyl
2-aminoisonicotinic acid in pyridine (45 mL) at 0 C
under an argon atmosphere was added acetyl chloride
dropwise over 5 min. After 2 h at 0 C, the reaction was
poured into over ice 300 g, extracted with ethyl acetate
35 (2 x 300 mL), washed with water (2 x 100 ml) followed by
brine (2 x 100 mL), and dried (Na₂SO₄). After
concentration in vacuo, the residue was purified by

application of flash chromatography (step gradient ethyl acetate: hexane 1:4 then ethyl acetate: hexane 1:1) to afford ethyl 2-acetamidoisonicotinate.

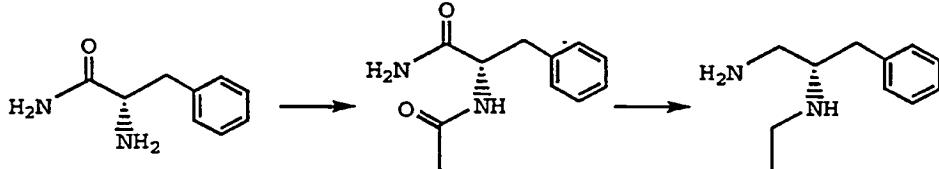
To a solution of diisopropylamine (14.15 mL, 101 mmol) and THF (40 mL) at -78 C was added n-butyl lithium (38.1 mL, 95 mmol) dropwise over 5 min. After 10 min, ethyl 4-fluorophenylacetate (17.3 g, 95 mmol) was added in 40 mL of dry THF. After 10 min, ethyl 2-acetamidoisonicotinate (6.0 g, 29 mmol) was added in 20 ml of dry THF. The reaction was allowed to warm to 23 C overnight, and then acetic acid (95 mmol) was added in one portion. The reaction was concentrated in vacuo, then partitioned repeatedly between saturated bicarbonate (200 ml) and ether (300 mL), the combined bicarbonate layers were neutralized with 10% citric acid, and extracted with ethyl acetate (2 x 300 mL). The organic layers were dried (Na₂SO₄), concentrated in vacuo to afford the Ethyl 2-(4-fluorophenyl)-3-oxo-3-(4-(2-acetamido)-pyridyl)-propionate.

Step B. 5-(4-fluorophenyl)-6-(4-(2-acetamido)pyridyl))-2-thiouracil:

Ethyl 2-(4-fluorophenyl)-3-oxo-3-(4-(2-acetamido)pyridyl)-propionate (1.3 g, 3.78 mmol) and thiourea (863 mg, 11.3 mmol) were suspended in anhydrous p-xylene (15 ml) with very efficient stirring. To the mixture pyridinium p-toluenesulfonate (38 mg) was added and refluxed for 12-16 h using a Dean-Stark apparatus with continuous removal of water (0.1 ml). Reaction mixture was cooled and a dark brown solid was filtered using a Buchner funnel. The collected solid was suspended in acetone (25 ml) and filtered. The acetone washed product contained a trace of thiourea, which was removed by trituration with hot water (20-30 ml). The product was filtered and air dried followed by azeotroping with toluene.

Example 21

Procedure for the preparation of (S)-2-N-Ethylamino-3-phenylpropylamine



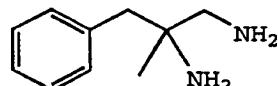
5 (S)-2-N-Ethylamino-3-phenylpropylamine: Acetic anhydride (1.2 ml, 12.7 mmol) was added to a stirring solution of L-phenylalanine amide (1.0 g, 6.10 mmol) in methanol (25 ml). After 1.5 h at room temperature, it was evaporated followed by drying in an oil pump vacuum.

10 The resultant L-N-ethylphenylalanine amide (6.1 mmol) was reduced with lithium aluminium hydride (570 mg, 15.0 mmol) in tetrahydrofuran (65 mmol) at 55°C for 4 h. The reaction mixture was poured into sat. aqu. sodium hydrogencarbonate followed by extraction with dichloromethane, drying and evaporation. Column chromatography on silica gel (chloroform : methanol : triethylamine = 90:7:3) provided the amine as a yellowish oil. MS (*m/z*): 179.1 (M+H)⁺; C₁₁H₁₈N₂ requir. 178.3.

20

Example 22

Procedure for the preparation of 2-Amino-2-methyl-3-phenylpropylamine

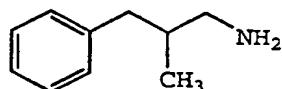


25 2-Amino-2-methyl-3-phenylpropylamine: A solution of commercially available D,L-a-methyl phenylalanine methyl ester (5.0 g, 25.7 mmol) in aqu. 28% ammonium hydroxide (50 ml) was kept at room temperature for 3 d. The resulting white precipitate of D,L-a-methyl phenylalanine amide was filtered and dried (2.5 g).

This material (2.0 g, 11.22 mmol) was reduced with lithium aluminium hydride (1.3 g, 34.26 mmol) in boiling tetrahydrofuran for 24 h. The reaction was quenched by the addition of sodium sulfate decahydrate at ice-bath 5 temperature. The salts were filtered off, followed by evaporation to leave the title compound as an oil. MS (m/z) : 165.1 ($M+H$)⁺; C₁₀H₁₆N₂ requir. 164.2. An alternative preparation was reported by M. Freiberger and R. B. Hasbrouck, J. Am. Chem. Soc. 82, 696-698 10 (1960).

Example 23

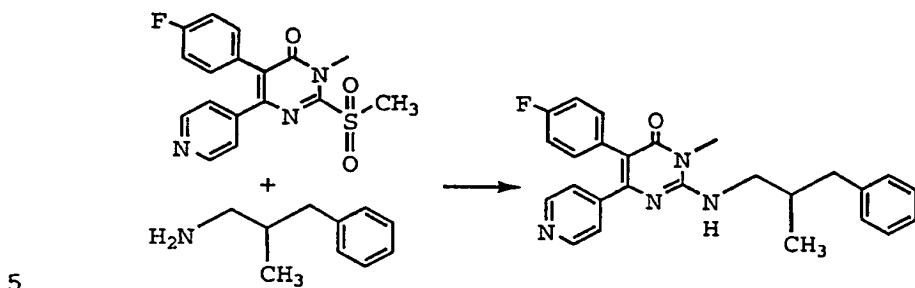
Procedure for the preparation of 2-Methyl-3-phenylpropylamine



15 2-Methyl-3-phenylpropylamine: A mixture of commercially available 2-methyl-3-phenylpropylamide (4.32 g, 26.5 mmol) and lithium aluminium hydride (1.3 g, 34.3 mmol) in tetrahydrofuran (184 ml) was stirred at room temperature for 5 h. It was poured into aqu. sat. 20 sodium sulfate and extracted with dichloromethane followed by drying of the organic solution and evaporation to provide the amine as an oil. Other syntheses have been reported, e.g. Dornow and Fust, Chem. Ber. 87, 984 (1954).

Example 24

Procedure for the preparation of 5-(4-Fluorophenyl)-3-methyl-2-((2-methy-3-phenylpropyl) amino)-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride

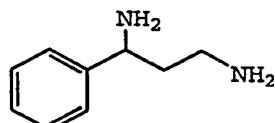


5-(4-Fluorophenyl)-3-methyl-2-((2-methy-3-phenylpropyl) amino)-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride:

A mixture of crude 5-(4-fluorophenyl)-3-methyl-2-methylsulfonyl-6-(4-pyridyl)-4(3H)-pyrimidinone (520 mg, 1.45 mmol) and 2-methyl-3-phenylpropylamine (1.5 g, 10.1 mmol) was heated at 50°C for 30 min. Column chromatography on silica gel (2-5% methanol/dichloromethane; hexane-acetone = 2 : 1) provided the title compound. MS (*m/z*) : 429.4 (M+H)⁺; C₂₆H₂₅FN₄O requir. 428.5 (free base).

Example 25

Procedure for the preparation of 1-Phenyl-1,3-propanediamine



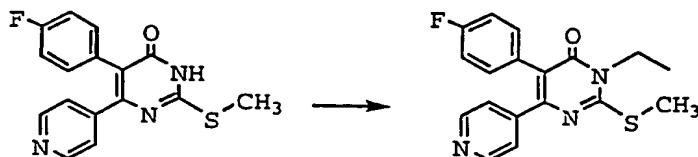
1-Phenyl-1,3-propanediamine: 3-Phenyl-3-aminopropionic acid (S. G. Cohen and S. Y. Weinstein, J. Am. Chem. Soc. 86, 725-728, 1964) was converted into 1-phenyl-1,3-propanediamine as reported in the literature (M. Kojima and J. Fujita, Bull. Chem. Soc. Jpn. 55, 1454-1459 (1982)).

145

Analogously, 1-(2-fluorophenyl)-1,3-propanediamine, 1-(2-methylphenyl)-1,3-propanediamine and 1-(2-chlorophenyl)-1,3-propanediamine have been prepared.

Example 26

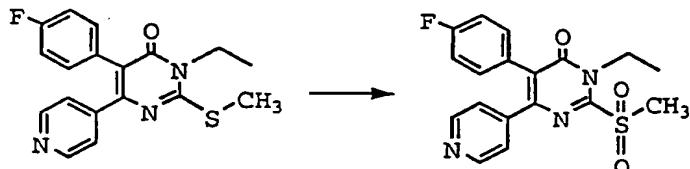
5 Procedure for the preparation of *3-Ethyl-5-(4-fluorophenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone*



10 *3-Ethyl-5-(4-fluorophenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone*: Ethyl bromide (600 ml, 8.03 mmol) was added to a stirred mixture of 5-(4-fluorophenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone (1.8 g, 5.97 mmol) and sodium hydride (60% oily suspension, 320 mg, 8 mmol) in *N,N*-dimethylformamide (60 ml) at room temperature. More ethyl bromide (2x 600 ml, 2x8.03 mmol) was added after 2 and 3.5 h. After 8 h, the reaction mixture was neutralized with acetic acid and evaporated. The remainder was taken up in dichloromethane, the organic solution was washed with water, dried and evaporated. Flash chromatography on a column of silica gel (hexane-acetone = 3:1, 2:1). provided in the second main fraction the title compound as a solid.

Example 27

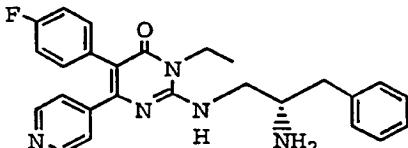
25 Procedure for the preparation of *3-Ethyl-5-(4-fluorophenyl)-2-methylsulfonyl-6-(4-pyridyl)-4(3H)-pyrimidinone*



3-Ethyl-5-(4-fluorophenyl)-2-methylsulfonyl-6-(4-pyridyl)-4(3H)-pyrimidinone: A mixture of 3-ethyl-5-(4-fluorophenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone (300 mg, 0.88 mmol) and Oxone® (potassium peroxyomonosulfate, 2.54 g, 4.14 mmol) in methanol (71 ml) and water (33 ml) was stirred for 14 h. The solvent was concentrated to about 35 ml, followed by extraction with dichloromethane, drying and evaporation. The resulting white solid was used without purification in the next step.

Example 28

Procedure for the preparation of 2-(((S)-2-Amino-3-phenylpropyl)-amino)-3-ethyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride



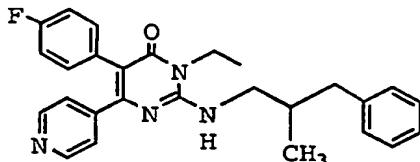
15

2-(((S)-2-Amino-3-phenylpropyl)-amino)-3-ethyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: A mixture of 3-ethyl-5-(4-fluorophenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidinone (150 mg, 0.44 mmol) and (S)-1,2-benzylethylendiamine (200 ml, ~1.3 mmol) was heated at 190°C for 4.5 h. Column chromatography on Iatrobeads® (chloroform : methanol : triethylamine = 90 : 7 : 3) provided the title compound as a free base which was converted into the crystallizing monohydrochloride by the addition of 2N hydrochloric acid (165 ml, 0.33 mmol) and methanol (1.5 ml). Filtration provided the title compound. MS (*m/z*) : 444.0 ($M+H$)⁺; C₂₆H₂₇FN₅O requir. 443.5 (free base).

30

Example 29

Procedure for the preparation of 3-Ethyl-5-(4-fluorophenyl)-2-((2-methy-3-phenylpropyl) amino)-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride



5

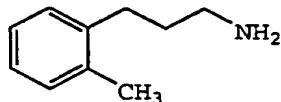
3-Ethyl-5-(4-fluorophenyl)-2-((2-methy-3-phenylpropyl) amino)-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride:

A mixture of crude 3-ethyl-5-(4-fluorophenyl)-2-methylsulfonyl-6-(4-pyridyl)-4(3H)-pyrimidinone (320 mg, 0.89 mmol) and 2-methyl-3-phenylpropylamine (600 ml, ~4 mmol) was heated at 60°C for 2 h. Column chromatography on silica gel (hexane-acetone= 2 : 1; 2-5% methanol/dichloromethane) provided the title compound. MS (*m/z*): 443.2 (M+H)⁺; C₂₇H₂₇FN₄O requir.

15 442.5.

Example 30

Procedure for the preparation of 3-(2-Methylphenyl)propylamine

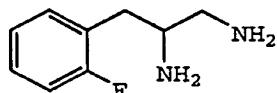


20 3-(2-Methylphenyl)propylamine: Diethyl cyanomethylphosphonate (5.0 ml, 30.9 mmol) was added to a stirring suspension of sodium hydride (60% oily suspension, 1.24 g, 31 mmol) in tetrahydrofuran (50 ml) under argon. After 30 min, 2-methylbenzaldehyde (3.6 ml, 31.1 mmol) was added and stirring continued for 1 h. The reaction was quenched by the addition of water and extracted with dichloromethane followed by drying and evaporation of the organic solution. Column chromatography (hexane; hexane : ethylacetate = 3 : 1)

provided 2-(2-methylphenyl)acrylonitrile as an oil. This material (3.8 g), 10% palladium on carbon (3.8 g) and 12 N hydrochloric acid (11.8 ml, 142 mmol) in methanol (125 ml) were hydrogenated with hydrogen at atmospheric pressure for 2 d. The catalyst was removed by filtration and the solvent was evaporated. The resultant material was partitioned between dichloromethane and water. The aqueous layer was made basic with 10 N sodium hydroxide and extracted with dichloromethane, followed by drying and evaporation. The resultant material was purified on a silica gel column (chloroform : methanol : triethylamine = 85 : 10 : 5) to provide the title compound as an oil.

Example 31

15 *Procedure for the preparation of 2-amino-3-(2-fluorophenyl)-propylamine*



Step A. Methyl 2-amino-3-(2-fluorophenyl)propionate: 5g (27.3 mmol) of (D,L)-(2-fluoro-phenyl)alanine was suspended in 50 ml methanolic HCl and stirred at room temperature for 3 days. The reaction mixture was concentrated in vacuo and dried to give a yellow oil. MS (m/z): 198 (M+H)⁺; C₁₀H₁₁FNO₂ requir. 197.2.

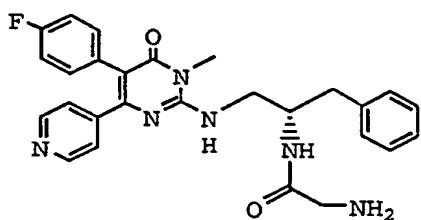
Step B. 2-Amino-3-(2-fluorophenyl)propionamide: Methyl 2-amino-3-(2-fluorophenyl) propionate was suspended in 50 ml 30% ammonium hydroxide and stirred at room temperature for 18 hrs. The mixture was filtered, washed with cold water and 2-amino-3-(2-fluorophenyl) propionamide was collected as a white solid. MS (m/z): 183.1 (M+H)⁺; C₉H₁₁FN₂O requir. 182.2.

Step C. 2-Amino-3-(2-fluorophenyl)-propylamine: 2-Amino-3-(2-fluorophenyl)propionamide was added carefully to a chilled (5°) mixture of LAH (1.0g, 26.3 mmol) and

20 ml THF under argon. The reaction was then heated at reflux for 10 hrs. The reaction was cooled to 5°C and carefully treated with Na₂SO₄•10 H₂O. The resulting mixture was stirred for 18 hrs, then filtered to remove 5 the solids. The filtrate was concentrated in vacuo to give an amber oil. MS (m/z): 169 (M+H)⁺; C₉H₁₁FN₂ requir. 168.19

Example 32

Procedure for the preparation of 5-(4-Fluorophenyl)-2-((S)-2-N-glycylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride



5-(4-Fluorophenyl)-2-((S)-2-N-glycylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: Ethyl chloroformate (56.8 µl, 0.59 mmol) was added at ice-bath temperature to a stirring mixture of N-(tert.-butoxycarbonyl)glycine (104 mg, 0.59 mmol) and 4-methylmorpholine (65.3 µl, 0.59 mmol) in tetrahydrofuran (9 ml). After 50 min, a solution of 2-((S)-2-amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone (250 mg, 0.58 mmol) in tetrahydrofuran (9 ml) was added at ice-bath temperature. Within 2 h, the mixture was allowed to reach room temperature. It was diluted with dichloromethane, washed with aqueous sodium hydrogencarbonate, followed by drying of the organic solution and evaporation. The resulting material was dissolved in methanol (1.2 ml) and 4N hydrogen chloride/dioxane (1.2 ml) was added. After 1 h at room temperature, it was evaporated and the remainder taken up in dichloromethane followed by washing with aqueous

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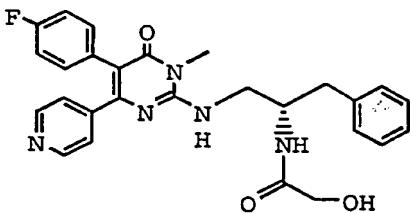
sodium hydrogencarbonate, drying of the organic solution and evaporation. Column chromatography on silica gel (dichloromethane - methanol - conc. ammonium hydroxide = 93 : 7 : 0.7) provided the title compound as the free base which was converted into the hydrochloride by the addition of 4N hydrogen chloride/dioxane (112 μ l, 0.45 mmol) to its methanolic solution (3 ml) followed by evaporation. MS (*m/z*): 487.1 ($M+H$)⁺; $C_{27}H_{27}FN_6O_2$ requir. 486.6 (free base).

10 Accordingly, 2-(((S)-2-N-glycylamino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride was prepared from 2-(((S)-2-amino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone.

15

Example 33

Procedure for the preparation of 5-(4-Fluorophenyl)-2-((S)-2-hydroxyacetamido-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone



20 5-(4-Fluorophenyl)-2-((S)-2-hydroxyacetamido-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: Acetoxyacetyl chloride (55 μ l, 0.51 mmol) was added at ice-bath temperature to a stirring solution of 2-(((S)-2-amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone (200 mg, 0.466 mmol) and triethylamine (130 μ l, 0.93 mmol) in dichloromethane (4 ml). After 50 min, the reaction was quenched by the addition of a drop of methanol followed by evaporation. The resultant material was taken up in a 30 1:1:1 mixture of methanol/water/triethylamine (3 ml) and

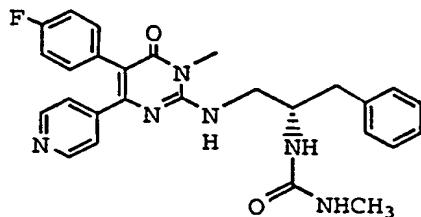
151

left overnight. Evaporation and subsequent column chromatography (3-7% methanol/chloroform) provided the title compound. MS (*m/z*): 488.3 (M+H)⁺; C₂₄H₂₆FN₆O, requir. 487.5.

5

Example 34

Procedure for the preparation of 5-(4-fluorophenyl)-2-(2-((3-N-methylureido)-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone



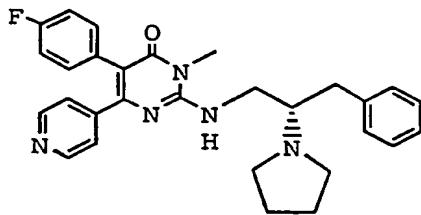
10 5-(4-Fluorophenyl)-2-(2-((3-N-methylureido)-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: Methyl isocyanate (6 μ l, 0.102 mmol) was added to a solution of 2-((S)-2-amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone (43.6 mg, 0.102 mmol) in dioxane (1.5 ml) at 15°C. After 15 min, the solvent was evaporated and the reaction product applied to a silica gel column (5-7% methanol/chloroform) to provide the title compound. MS (*m/z*): 486.6 (M+H)⁺; C₂₄H₂₆FN₆O₂, requir. 486.6.

15

20

Example 35

Procedure for the preparation of 5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-2-((2-pyrrolidinyl-3-phenylpropyl)-amino)-4(3H)-pyrimidinone hydrochloride



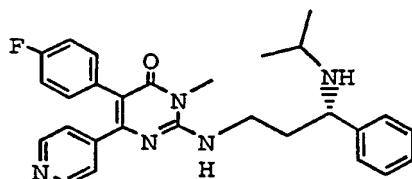
5-(4-Fluorophenyl)-3-methyl-6-(4-pyridyl)-2-((S)-2-pyrrolidinyl-3-phenylpropyl)-amino)-4(3H)-pyrimidinone hydrochloride:

Sodium hydride (60% oily suspension, 84 mg, 2.1 mmol) was added to a solution of 2-((S)-2-amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone (300 mg, 0.70 mmol) in *N,N*-dimethylformamide (8 ml) at ice-bath temperature. After 30 min, 1,4-dibromobutane (108 µl, 0.91 mmol) was added. Stirring was continued for 30 min at ice-bath temperature, then 20 h at room temperature. It was neutralized with acetic acid, followed by evaporation. The crude product was purified on a column of silica gel (dichloromethane - methanol = 93 : 7; dichloromethane - methanol - conc. ammonium hydroxide = 93 : 7 : 0.7). The resultant product was converted into the hydrochloride by the addition of 4N hydrogen chloride/dioxane (37 µl) to its methanolic solution (2 ml) and subsequent evaporation. MS (*m/z*): 484.6 (M+H)⁺; C₂₉H₃₀FN₃O requir. 483.6 (free base).

20

Example 36

Procedure for the preparation of 5-(4-fluorophenyl)-2-((S)-3-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride



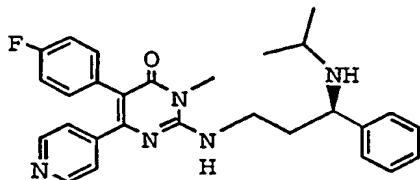
25

5-(4-Fluorophenyl)-2-((S)-3-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: Sodium triacetoxyborohydride (12.9 mg, 0.061 mmol) was added to a stirring mixture of 2-((S)-3-amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone (21.8 mg, 0.051 mmol) and acetone (4.5 µl, 0.061 mmol)

in 1,2-dichloroethane (0.4 ml). After 2.5 h, the reaction was quenched by the addition of sat. aqu. sodium hydrogencarbonate, followed by extraction with dichloromethane, drying of the organic solution and evaporation. Chromatography on a column of silica gel (10% methanol/chloroform) provided the title compound as a free base which was converted into the monohydrochloride by the addition of 4N hydrochloric acid/dioxane (12.2 μ l) to its methanolic solution (1 ml) and subsequent evaporation. MS (*m/z*): 472.0 ($M+H$) $^+$; C₂₈H₃₀FN₅O requir. 471.6 (free base).

Example 37

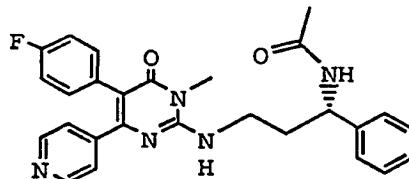
*Procedure for the preparation of 5-(4-fluorophenyl)-2-((*R*)-3-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3*H*)-pyrimidinone hydrochloride*



5-(4-Fluorophenyl)-2-((*R*)-3-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3*H*)-pyrimidinone hydrochloride was prepared from 5-(4-fluorophenyl)-2-((*R*)-3-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3*H*)-pyrimidinone as described above for its S-enantiomer. MS (*m/z*): 472.1 ($M+H$) $^+$; C₂₈H₃₀FN₅O requir. 471.6 (free base).

Example 38

*Procedure for the preparation of 2-((*S*)-3-acetamido-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3*H*)-pyrimidinone*



2-(((S)-3-Acetamido-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone: A solution of 2-(((S)-3-amino-3-

5 phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone (23.8 mg, 0.055 mmol) and acetic anhydride (20 μ l, 0.21 mmol) in methanol (1 ml) was kept for 30 min at room temperature. Evaporation was followed by column chromatography (dichloromethane -
10 methanol - ammonium hydroxide = 93 : 7 : 0.7) to provide the title compound. MS (*m/z*): 472.2 ($M+H$)⁺; C₂₇H₂₆FN₅O₂ requir. 471.5.

Example 39

Procedure for the preparation of (S)-1-Phenyl-1,3-propanediamine



(S)-1-Phenyl-1,3-propanediamine: S-3-N-tert--

Butoxycarbonylamino-3-phenylpropionitrile was prepared according to the literature (W.J. Wheeler and D.D.

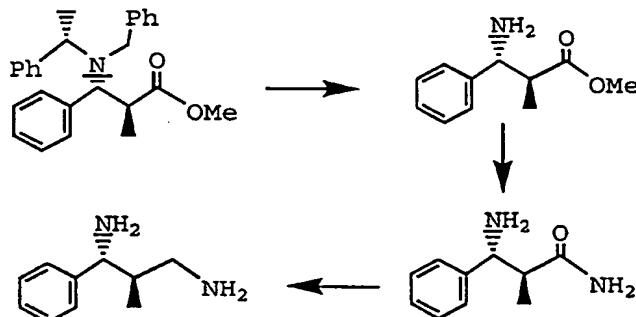
20 O'Bannon, J. Label. Compds. Radiopharm. **XXXI** (4), 305-315, 1992) from D-(-)- α -phenylglycinol. For reduction (D. Mitchell and T.M. Koenig, Synth. Comm. **25** (8), 1231-1238, 1995), borane-methyl sulfide complex (2N, 3 ml, 6 mmol) was added dropwise to a solution of the nitrile (1 g, 4.06 mmol) in tetrahydrofuran (6 ml). Methyl sulfide was distilled off and the resulting solution refluxed for 2.5 h. With ice-cooling, methanolic hydrogen chloride (1N, 3 ml) was added followed by evaporation.

The remainder was taken up in methanol (10 ml) and 4N hydrogen chloride/dioxane (10 ml) was added. After 1 h at room temperature, it was evaporated and the aqueous solution of the resultant product was washed with dichloromethane. The aqueous solution was made basic by the addition of solid potassium hydroxide followed by repeated dichloromethane extractions. Drying and evaporation of the dichloromethane solution left the crude diamine as an oil. MS (*m/z*): 150.8 (M+H)⁺; C₉H₁₄N₂ requir. 150.2.

Enantiomeric *(R)*-1-phenyl-1,3-propanediamine was prepared analogously from L-(+)- α -phenylglycinol. MS (*m/z*): 150.9 (M+H)⁺; C₉H₁₄N₂ requir. 150.2.

Example 40

15 Procedure for the preparation of (2*R*,3*R*)-2-methyl-3-phenyl-1,3-propanediamine



Step A: Methyl (2*S*,3*R*, α *S*)-3-(*N*-benzyl-*N*- α -methylbenzylamino)-2-methyl-3-phenylpropionate was prepared as reported for the 2*R*,3*S*, α *R*-enantiomer (S.G. Davies and I.A.S. Walters, J. Chem. Soc. Perkin Trans.I, 1129-1139 (1994)).

Step B: Methyl (2*S*,3*R*)-3-amino-2-methyl-3-phenylpropionate: A mixture of methyl (2*S*,3*R*, α *S*)-3-(*N*-benzyl-*N*- α -methylbenzylamino)-2-methyl-3-phenylpropionate (13.0 g, 33.55 mmol) and 10% palladium-

on-carbon (13.0 g) in glacial acetic acid (260 ml) was hydrogenated under a balloon of hydrogen for 24 h. The catalyst was removed by filtration followed by evaporation and co-distillation with toluene to provide 5 the title compound as a white solid. MS (*m/z*): 194.2 (*M+H*)⁺; C₁₁H₁₃NO requir. 193.3.

Step C: (2S,3R)-3-Amino-2-methyl-3-phenylpropionamide:

A solution of methyl (2S,3R)-3-amino-2-methyl-3-phenylpropionate (6.3 g, 33 mmol) in 2N methanolic 10 ammonia (20 ml) and ammonium hydroxide (28-30%, 40 ml) was stirred at room temperature. After 4d, it was evaporated followed by chromatography on a short column 15 of silica gel (dichloromethane - methanol - conc. ammonium hydroxide = 93 : 7 : 0.7; 90 : 10 : 0.8) to provide the amide as a white solid. MS (*m/z*): 179.2 (*M+H*)⁺; C₁₀H₁₄N₂O requir. 178.2.

Step D: (2R,3R)-2-methyl-3-phenyl-1,3-propanediamine:

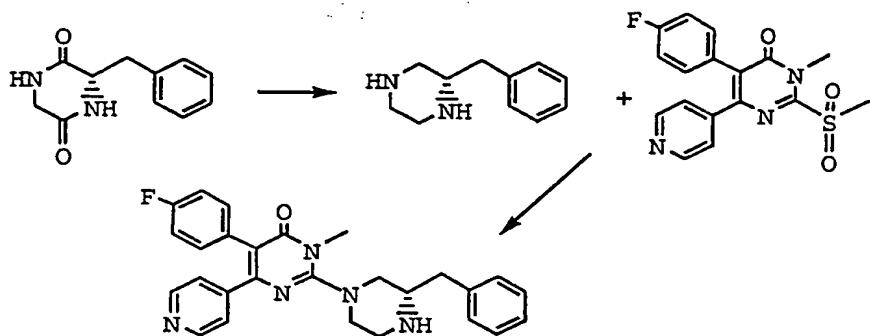
Lithium aluminium hydride (2.3 g, 60.60 mmol) was added 20 in portions to a stirring solution of (2S,3R)-3-amino-2-methyl-3-phenylpropionamide (2.6 g, 14.59 mmol) in tetrahydrofuran (54 ml) at ice-bath temperature. After 45 min, the mixture was heated at reflux for 16 h. With ice-bath cooling, the reaction was quenched by the portionwise addition of sodium sulfate decahydrate and 25 some methanol until hydrogen evolution ceased. The solids were removed by filtration and washed with dichloromethane. The combined filtrates were evaporated to provide the title compound. MS (*m/z*): 165.2 (*M+H*)⁺; C₁₀H₁₄N₂ requir. 164.3.

30 Accordingly, the enantiomer (2S,3S)-2-methyl-3-phenyl-1,3-propanediamine was prepared from methyl (2R,3S, α R)-3-(*N*-benzyl-*N*- α -methylbenzylamino)-2-methyl-3-phenylpropionate. MS (*m/z*): 165.3 (*M+H*)⁺; C₁₀H₁₄N₂ requir. 164.3.

Analogously, the enantiomers (2R,3S)-2-methyl-3-phenyl-1,3-propanediamine and (2S,3R)-2-methyl-3-phenyl-1,3-propanediamine may be prepared from tert.butyl (2S,3S,αR)- and -(2R,3R,αS)-3-(N-benzyl-N-α-methylbenzylamino)-2-methyl-3-phenylpropionate (S. Davies et al., J. Chem. Soc. Chem. Commun. 1153-1155, 1993).

Example 41

Procedure for the preparation of 2-((S)-3-Benzylpiperaziny)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride



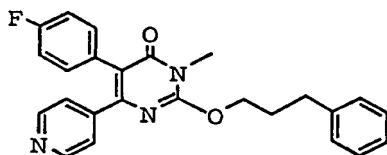
Step A: (S)-2-Benzylpiperazine: At ice-bath temperature, lithium aluminium hydride (1.6 g, 42.16 mmol) was added in portions to a stirring mixture of (S)-2-benzylpiperazine-3,6-dione (3.0 g, 14.70 mmol) (comm. avail.) and tetrahydrofuran (80 ml). After 30 min at ice-bath temperature, the mixture was refluxed for 4 h with stirring. The reaction was quenched by the portionwise addition of sodium sulfate decahydrate and some methanol until hydrogen evolution ceased. It was filtered and the solids were washed several times with dichloromethane. The combined filtrates were evaporated to leave a white solid. MS (*m/z*): 177.1 ($M+H$)⁺; C₁₁H₁₆N₂ requir. 176.3.

Step B: 2-((S)-3-Benzylpiperaziny)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride:

A mixture of crude 5-(4-fluorophenyl)-3-methyl-2-methylsulfonyl-6-(4-pyridyl)-4(3H)-pyrimidinone (434 mg, 1.21 mmol) and (S)-2-benzylpiperazine (426 mg, 2.42 mmol) was heated at 105°C for 1 h. The crude reaction product was purified by column chromatography on silica gel (dichloromethane - methane = 93 : 7; dichloromethane - methanol - conc. ammonium hydroxide = 93 : 7: 0.7). The resulting material was converted into its hydrochloride by the addition of 4N hydrogen chloride/dioxane (75 µl) to its methanolic solution (3 ml) followed by evaporation. MS (*m/z*): 456.5 (M+H)⁺; C₂₇H₂₆FN₃O requir. 455.5 (free base).

Example 42

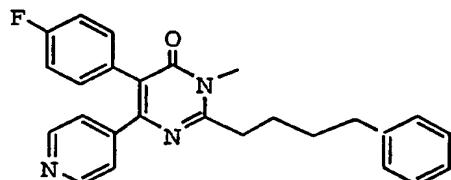
Procedure for the preparation of 5-(4-fluorophenyl)-3-methyl-2-(3-phenylpropoxy)-6-(4-pyridyl)-4(3H)-pyrimidinone



5-(4-fluorophenyl)-3-methyl-2-(3-phenylpropoxy)-6-(4-pyridyl)-4(3H)-pyrimidinone: Sodium hydride (60% oily suspension, 111 mg, 2.79 mmol) was added to a stirred solution of 3-phenylpropanol (387 mg, 2.85 mmol) in tetrahydrofuran (1 ml). After gas evolution ceased, 5-(4-fluorophenyl)-3-methyl-2-methylsulfonyl-6-(4-pyridyl)-4(3H)-pyrimidinone (100 mg, 0.279 mmol) was added and the mixture was heated at 60°C for 30 min. The reaction mixture was partitioned between dichloromethane and water. The organic solution was washed with brine, dried and evaporated. Column chromatography on silica gel (hexane - ethyl acetate = 2 : 1) provided the title compound. MS (*m/z*): 416.1 (M+H)⁺; C₂₅H₂₂FN₃O₂ requir. 415.5.

Example 43

Procedure for the preparation of 5-(4-fluorophenyl)-3-methyl-2-(4-phenylbutyl)-6-(4-pyridyl)-4(3H)-pyrimidinone



5

Step A: 5-(4-Fluorophenyl)-2-(4-phenylbutyl)-6-(4-pyridyl)-4(3H)-pyrimidinone: Ethyl 2-(4-fluorophenyl)-3-oxo-3-(4-pyridyl)-propionate (293 mg, 1.02 mmol), 4-phenylbutanecarboxamidine (315 mg, 1.79 mmol) and

10 pyridinium p-toluenesulfonate (10 mg) were suspended in p-xylene (10 ml). With efficient stirring, the mixture was heated to reflux using a Dean-Stark apparatus with continuous removal of water. After 16 h, the solvent was evaporated and the product purified by column chromatography on silica gel (3% methanol/dichloromethane) followed by recrystallization from acetone. MS (*m/z*): 400.3 (M+H)⁺; C₂₅H₂₂FN₃O requir. 399.5.

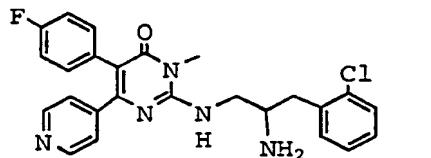
Step B: 5-(4-Fluorophenyl)-3-methyl-2-(4-phenylbutyl)-6-(4-pyridyl)-4(3H)-pyrimidinone:

20 Methyl iodide (22 µl, 0.351 mmol) was added to a stirring mixture of 5-(4-fluorophenyl)-2-(4-phenylbutyl)-6-(4-pyridyl)-4(3H)-pyrimidinone (140 mg, 0.351 mmol) and potassium carbonate (49 mg, 0.351 mmol) in *N,N*-dimethylformamide (5 ml). After 75 min, it was evaporated and the resultant product purified on a silica gel column (hexane - acetone = 3 : 1; 2 : 1) to provide the title compound. MS (*m/z*): 414.3 (M+H)⁺; C₂₆H₂₄FN₃O requir. 413.5.

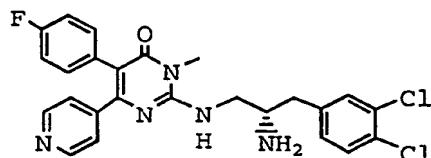
30

Example 44

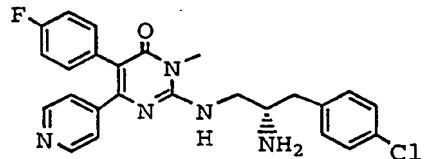
The compounds shown in Table I were prepared using the procedures of Examples 1-43.

TABLE I

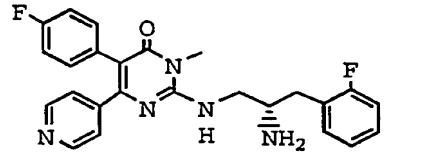
MS (*m/z*): 464.0 (M)⁺;
C₂₅H₂₃FN₅O requir. 463.9



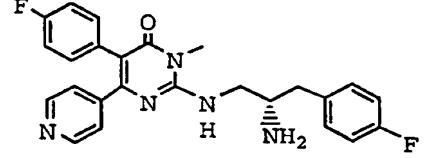
MS (*m/z*): 498.0 (M)⁺;
C₂₅H₂₂FN₅O requir. 498.4



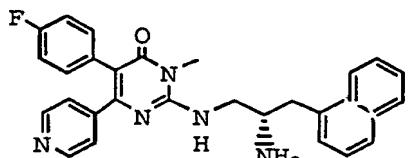
MS (*m/z*): 464.1 (M)⁺;
C₂₅H₂₃ClFN₅O requir. 463.9



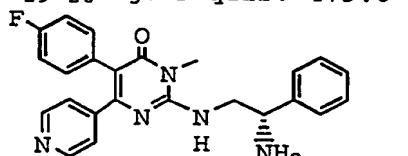
MS (*m/z*): 448.3 (M+H)⁺;
C₂₅H₂₃F₂N₅O₂ requir. 447.5



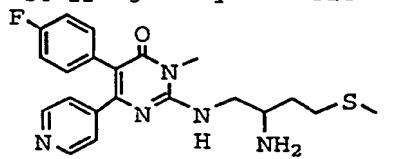
MS (*m/z*): 448.2 (M+H)⁺;
C₂₅H₂₂F₂N₅O requir. 447.3



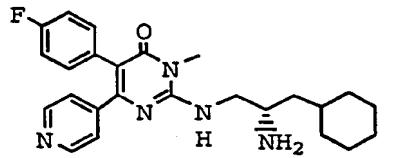
MS (*m/z*): 479.7 (M)⁺;
C₂₉H₂₆FN₅O requir. 479.6



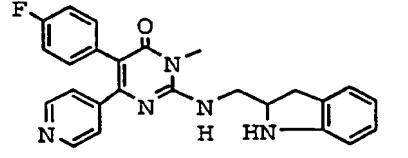
MS (*m/z*): 416.1 (M+H)⁺;
C₂₄H₂₂FN₅O requir. 415



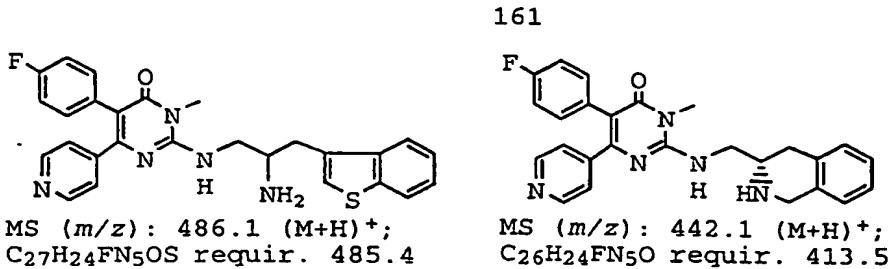
MS (*m/z*): 414.0 (M+H)⁺;
C₂₁H₂₄FN₅OS requir. 413.5



MS (*m/z*): 436.2 (M+H)⁺;
C₂₅H₃₀FN₅O requir. 435.6

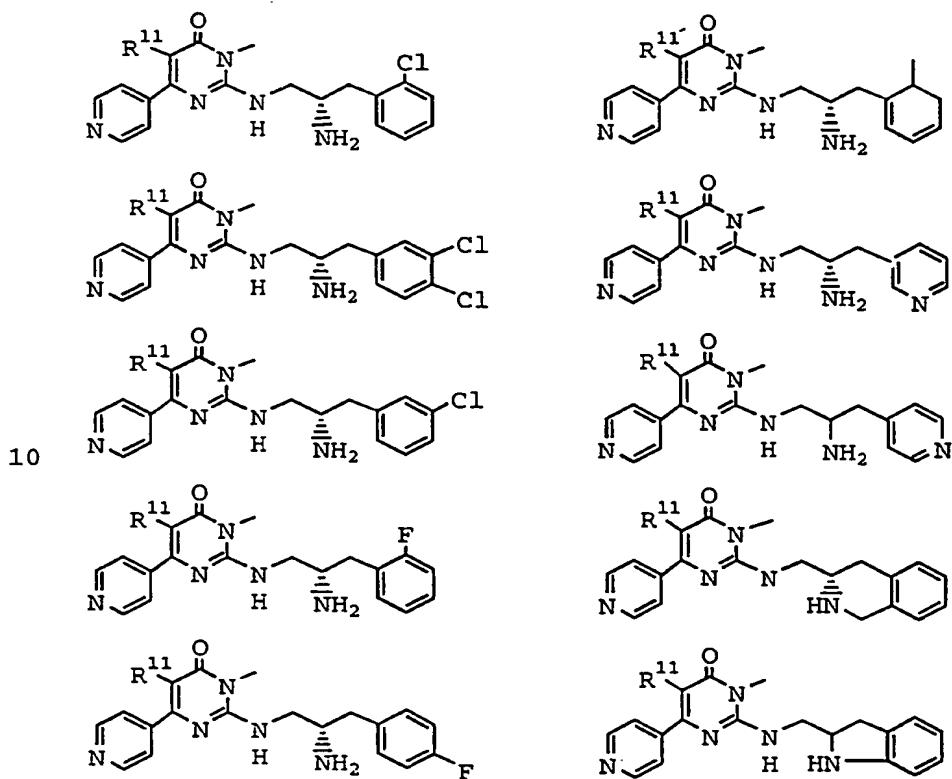


MS (*m/z*): 428.1 (M+H)⁺;
C₂₅H₂₂FN₅O requir. 427.5

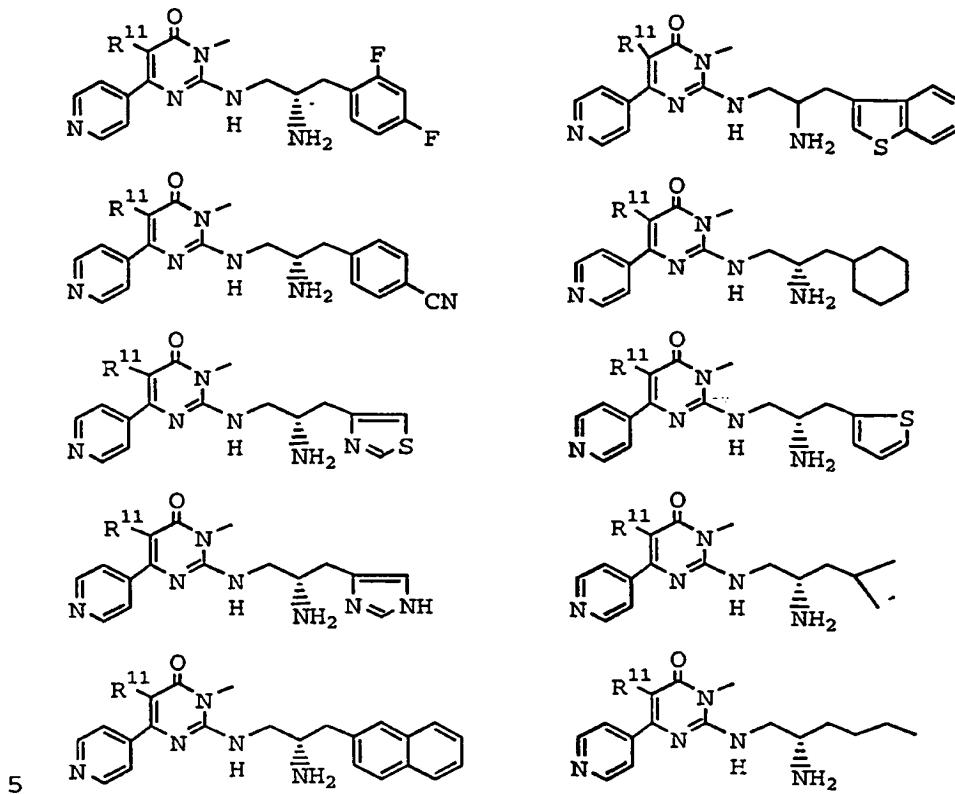
**Example 45**

The compounds shown in Table II can be prepared using the procedures of Examples 1-43, wherein R¹¹ represents 3-methylphenyl, 3-chlorophenyl, 3-trifluoromethylphenyl, 4-fluorophenyl, 4-methylphenyl, 4-chlorophenyl and 3,4-dimethylphenyl.

5

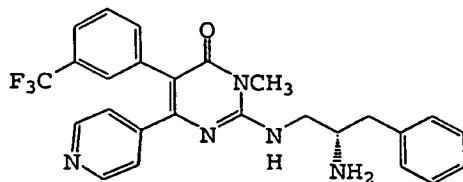
TABLE II

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**Example 46**

Procedure for the preparation of 3-methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-trifluoromethyl phenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

10



Step A. 6-(4-pyridyl)-2-thiouracil: Ethyl isonicotinoylacetate (5g, 25.89 mmol) and thiourea (5.94 g, 77.64 mmol) were suspended in anhydrous p-xylene (100ml) with vigorous stirring.

To the mixture, 15 pyridinium p-toluenesulfonate (150mg) was added and refluxed for 12-16 h using a Dean-Stark apparatus with

continuous removal of water (0.5ml). The reaction mixture was cooled and a dark brown solid was filtered. The collected solid was suspended in acetone (25 ml) and filtered. The acetone washed product contain trace of thiourea, which was removed by trituration with hot water (20-30ml). The title compound was isolated by filtration. MS (m/z): 206.2 C₁₁H₁₁N₃OS requir. 205.3. ¹H-NMR (DMSO-d₆): d 12.65 (bm, 2H, NH and SH), 8.71(m, 2H, pyrid.), 7.66(m, 2H, Pyrid.), 6.25 (s, 1H, H-5).

10 Step B. 3-Methyl-6-(4-pyridyl)-2-methylthio-4(3H)-pyrimidinone: 6-(4-Pyridyl)-2-thiouracil (1.5g 7.299 mmol) was dissolved in DMF (50 ml) and the mixture was cooled to 0°C. Sodium hydride (0.437 g, 0.730g 60% in oil, 18.25 mmol) was added and the reaction mixture was stirred for 30 minutes. Methyl iodide (1.2 ml, 2.6g, 18.25 mmol) was added dropwise over 15 minutes. Formation of dimethyl compound was monitored by TLC. Reaction mixture was concentrated and the residue chromatographed on silica gel column using hexane: acetone (9:1, 4:1 and 2:1) to obtain the title compound as a solid: MS(m/z):234.1 C₁₁H₁₁N₃OS requir. 233.2; ¹H-NMR(CDCl₃):d 8.75 (m, 2H, pyridyl), 7.8 (m, 2H, pyridyl), 6.75 (s, 1H), 3.58 (s, 3H, N-CH₃), 2.72 (s, 3H, S-CH₃).

20 Step C. 3-Methyl-5-bromo-6-(4-pyridyl)-2-methylthio-4(3H)-pyrimidinone: 3-Methyl-6-(4-pyridyl)-2-methylthio-4(3H)-pyrimidinone (1.00g 4.29 mmol) was dispersed in acetic acid (24 ml) and to the clear solution Bromine (0.5ml, 1.5g 9.38 mmol) was added. The reaction mixture was stirred at room temperature for 24 h. The mixture was concentrated and the residue was co-evaporated with toluene until all bromine is removed. The crude compound is ready to use in next step. MS(m/z): 312 and 314. C₁₁H₁₀BrN₃OS requir. 311 and 313. ¹H-NMR(DMSO-d₆):d 8.75 (m, 2H, pyridyl) 8.19 (m, 2H, pyridyl), 3.67 (s, 3H, N-CH₃), 2.80 (s, 3H, S-CH₃).

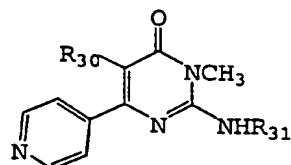
Step D. 3-Methyl-5-(3-trifluoromethylphenyl)-6-(4-pyridyl)-2-thiomethyl-4(3H)-pyrimidinone: 3-Methyl-5-bromo-6-(4-pyridyl)-2-methylthio-4(3H)-pyrimidinone (1.2g, 3.8 mmol) was dispersed in 2M sodium carbonate solution (30 ml) and the pale yellow colour of the adhered bromine disappeared to give colourless precipitate in the reaction mixture. 3-Trifluoromethylbenzene boronic acid (1.00 g, 5.27 mmol) and toluene (30ml) were added to the above mixture and the reaction mixture was degassed. Tetrakis triphenyl phosphine Pd(0) (350 mg) was added. The reaction mixture was refluxed for 8-12h. The formation of the product was monitored by TLC. The mixture was cooled, diluted with toluene(20ml) and washed with water. The organic layer was dried over sodium sulfate, concentrated and product isolated by silica gel chromatgraphy to give the titled compoud. MS(m/z): 378.4 C₁₈H₁₄F₃N₂OS requir. 377.39; 1H-NMR(CDCl₃):d 8.5 (m, 2H, pyridyl), 7.45 (s,1H), 7.17-7.25 (m, 3H, pyridyl and Ph-CF₃), 6.95 (d, 1H, Ph-CF₃), 3.67 (N-CH₃), 2.8 (S-CH₃).

Step E. 3-methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone: 3-Methyl-5-(3-trifluoromethylphenyl)-6-(4-pyridyl)-2-thiomethyl-4(3H)-pyrimidinone (0.7g, 1.85 mmol) and (S)-2-amino-3-phenyl-1-propylamine (0.9 ml, 6.00 mmol) were mixed in a round bottom flask and heated at 185°C for 3h. The mixture was separated on silica gel (dichloromethane: methanol: ammonium hydroxide 92:7:1) to obtain compound titled compound. MS(m/z): 480, C₂₆H₂₄F₃N₅O requir 479.51; 1H-NMR(CDCl₃):d 8.49 (m, 2H, pyridyl), 7.51-7.17 (m, 11H, Ph and pyridyl), 5.81 (bm, 1H, NH), 3.91 (m, 1H, CH), 3.53 (s, 3H, N-CH₃), 3.35 (m, 2H, CH₂), 2.94 (dd, 1H, CH₂), 2.82 (dd, 1H, CH₂).

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Example 47

Using the corresponding starting materials, the following compounds of Table III were prepared using the procedure for 3-methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone.

TABLE III

	<u>R₁₀</u>	<u>R₁₁</u>	<u>MS (m/z)</u>
10	4-tolyl 4-trifluoromethyl phenyl	2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl	426 480
15	3-isopropylphenyl 3-chloro-4-fluoro phenyl 3,5-bis(trifluoro methyl)phenyl	2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl	454 464
20	3,4-dichloro phenyl 1-naphthyl 3-fluorophenyl 3-chlorophenyl	2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl	482 462 430
25	3-methylphenyl 4-chlorophenyl 2-chlorophenyl 2-thienyl	2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl	427 440.6 467
30	3,4-dimethylphenyl 3,5-dichloro phenyl 4-tolyl 3-trifluoromethyl phenyl	2(S)-amino-3-phenyl-propyl 3-phenylpropyl 3-phenylpropyl	465 411
35	4-methoxyphenyl 4-trifluoromethyl phenyl 3-chlorophenyl 3-methylphenyl 4-chlorophenyl 2-chlorophenyl	3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl	465 427
40	3-nitrophenyl	3-phenylpropyl	465

	3-methoxyphenyl	3-phenyl-propyl	
	2-fluorophenyl	3-phenyl-propyl	
	benzothienyl	3-phenyl-propyl	
	3-fluorophenyl	2-methyl-3-phenyl-propyl	429
5	1-naphthyl	2-methyl-3-phenyl-propyl	461
	3-trifluoromethyl	2(S)-dimethylamino-	
	phenyl	3-phenylpropyl	
	3-methylphenyl	2(S)-dimethylamino-	
		3-phenylpropyl	
10	3-chlorophenyl	2(S)-N,N-dimethylamino-	
		3-phenylpropyl	
	3-nitrophenyl	2(S)-N,N-dimethylamino-	
		3-phenylpropyl	
	3-methoxyphenyl	2(S)-N,N-dimethylamino-	
15		3-phenylpropyl	
	2-fluorophenyl	2(S)-N,N-dimethylamino-	
		3-phenylpropyl	
	3-trifluoromethyl	(S)-tetrahydroisoquinol-3-	492.1
	phenyl	ylmethylenamino	
20	3-methylphenyl	(S)-tetrahydroisoquinol-3-	438
		ylmethylenamino	
	3,4-dimethylphenyl	3-amino-3-phenylpropylamine	440.6
	3-methylphenyl	3-amino-3-phenylpropylamine	
	benzothienyl	3-amino-3-phenylpropylamine	
25	benzofuranyl	3-amino-3-phenylpropylamine	

Example 48

3-Methyl-5-(4-methylsulfinylphenyl)-6-(4-pyridyl)-2-thiomethyl-4(3H)-pyrimidinone: The title compound was prepared in the manner of example 34-D substituting 4-methylsulfinylbenzene boronic acid for 3-trifluoromethylbenzene boronic.

Example 49

3-methyl-2-(3(S)-(1,2,3,4-tetrahydroisoquinolinyl)methylamino)-5-(4-methylthiophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone: The title compound was prepared in the manner of example 34 step D with the following substitutions of 3-methyl-5-(4-methylsulfinylphenyl)-6-(4-pyridyl)-2-thiomethyl-4(3H)-pyrimidinone for 3-methyl-5-(3-trifluoromethylphenyl)-6-(4-pyridyl)-2-thiomethyl-4(3H)-pyrimidinone and 3(S)-(1,2,3,4-tetrahydroisoquinolinyl)methylamine for (S)-2-amino-3-phenyl-1-propylamine: MS (m/z) 470 (M+H)+.

Example 50

3-methyl-2-(3(S)-(1,2,3,4-tetrahydroisocuinolinyl)methylamino)-5-(4-methylsulfonylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone: To a solution of 3-methyl-2-(3(S)-(1,2,3,4-tetrahydroisoquinolinyl)methylamino)-5-(4-methylthiophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone (50 mg, 0.11 mmol) in methanol:water (15 mL:10 mL) was added oxone (127 mg, 0.21 mmol) as a solid in one portion at 23°C. After 16 h, the reaction was concentrated under a stream of nitrogen. The reaction mixture was applied directly to purification via preparative plate chromatography (3 silica gel 2mm thick plates; 5% methanol in methylene chloride) to afford the title compound : MS (m/z) 502 (M+H)⁺.

15 **Example 51**

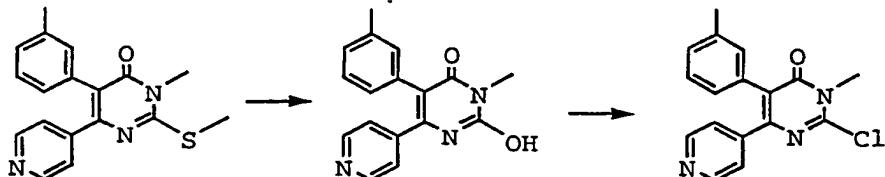
2-(((S)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-5-(3-trifluoromethylphenyl)-4(3H)-pyrimidinone hydrochloride was prepared from 3-methyl-2-methylthio-6-(4-pyridyl)-5-(3-trifluoromethylphenyl)-4(3H)-pyrimidinone and (S)-1-phenyl-1,3-propanediamine according to the General Procedure. The reaction was at 190°C for 1 h. MS (m/z) : 480.0 (M+H)⁺; C₂₆H₂₄F₃N₅O requir. 479.5 (free base).

Example 52

25 2-(((R)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-5-(3-trifluoromethylphenyl)-4(3H)-pyrimidinone hydrochloride was prepared from 3-methyl-2-methylthio-6-(4-pyridyl)-5-(3-trifluoromethylphenyl)-4(3H)-pyrimidinone and (R)-1-phenyl-1,3-propanediamine according to the General Procedure. The reaction was done at 190°C for 3.5 h. MS (m/z) : 480.4 (M+H)⁺; C₂₆H₂₄F₃N₅O requir. 479.5 (free base).

Example 53

Procedure for the preparation of 2-chloro-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone



5 Step A: 3-Methyl-5-(3-methylphenyl)-6-(4-pyridyl)-2,4(1H,3H)-pyrimidindione: 10 N Sodium hydroxide (25 ml) and water (50 ml) was added to a solution of 3-methyl-5-(3-methylphenyl)-2-methylthio-6-(4-pyridyl)-4(3H)-pyrimidindione (16.17 g, 0.05 mol) in dioxane (65 ml). The mixture was heated at 80°C for 16 h under argon. The mixture was allowed to reach room temperature and the pH value was adjusted to 9 with 1 N hydrochloric acid. The precipitate was filtered, washed with water and dried to give the title compound. MS
10 (m/z): 292 (M-H)⁺; C₁₇H₁₅N₃O, requir. 293.3.

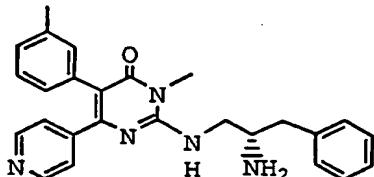
15 Step B: 2-Chloro-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone: A mixture of 3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-2,4(1H,3H)-pyrimidindione (12.5 g, 0.043 mol) and phosphorus oxychloride (65 ml) was refluxed for 16 h. The excess of phosphorus oxychloride was evaporated followed by co-distillation with toluene. The remainder was carefully partitioned between dichloromethane and aqueous sodium hydrogencarbonate. The organic solution was washed with water, dried and evaporated to leave the title compound.
20 MS (m/z): 312 (M)⁺; C₁₇H₁₄ClN₃O requir. 311.8.

25 2-Chloro-3-methyl-6-(4-pyridyl)-5-(3-trifluoromethylphenyl)-4(3H)-pyrimidinone was prepared according to the same procedure.

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Example 54

Procedure for the preparation of 2-((S)-2-amino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride



5

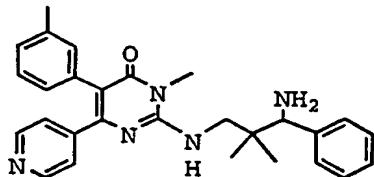
2-((S)-2-Amino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

hydrochloride: A solution of 2-chloro-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone (3.34 g, 10.71 mmol) and (S)-1-benzyl-1,2-ethanediamine (2.3 g, 15.31 mmol) in ethanol (50 ml) was stirred at room temperature for 16 h. The solvent was evaporated and the crude product recrystallized from methanol. MS (m/z): 426 (M+H)⁺; C₂₆H₂₇N₅O requir. 425.5 (free base).

15

Example 55

Procedure for the preparation of 2-((3-amino-2,2-dimethyl-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride



20

2-((3-Amino-2,2-dimethyl-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone hydrochloride: A solution of 2-chloro-3-

25 methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone (228 mg, 0.73 mmol) and 3-phenyl-2,2-dimethyl-1,3-propanediamine (178 mg, 1 mmol) (prepared according to: W. Ten Hoeve and H. Wynberg, *Synth. Commun.*

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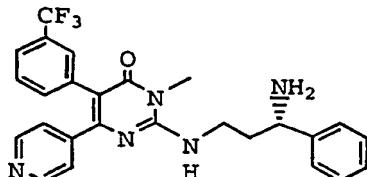
24 (15), 2215-2221, 1994) in ethanol (4 ml) was stirred at room temperature for 16 h. The solvent was evaporated and the crude product purified by column chromatography on silica gel. MS (*m/z*): 454 (M+H)⁺; 5 C₂₈H₃₁N₅O requir. 453.6 (free base).

Accordingly, 2-((-3-Amino-2,2-dimethyl-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-5-(3-trifluoromethylphenyl)-4(3H)-pyrimidinone hydrochloride was prepared. MS (*m/z*): 508 (M+H)⁺; C₂₈H₂₈F₃N₅O requir. 507.6 (free base).

10

Example 56

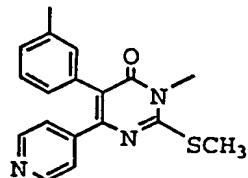
Procedure for the preparation of 2-(((S)-3-amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-5-(3-trifluoromethylphenyl)-4(3H)-pyrimidinone hydrochloride



15 2-(((S)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-5-(3-trifluoromethylphenyl)-4(3H)-pyrimidinone hydrochloride: Aqueous sat. sodium carbonate (2 ml) was added to a solution of 2-chloro-3-methyl-6-(4-pyridyl)-5-(3-trifluoromethylphenyl)-4(3H)-pyrimidinone hydrochloride (730 mg, 2 mmol) and (S)-1-phenyl-1,3-propanediamine (360 mg, 2.4 mmol) in ethanol (10 ml). The mixture was stirred for 4 h at room temperature. It was evaporated and the remainder partitioned between dichloromethane and water. The organic solution was dried and evaporated followed by column chromatography on silica gel (dichloromethane : methanol : conc. ammonium hydroxide = 93 : 7 : 0.7). MS (*m/z*): 480 (M+H)⁺; 20 C₂₆H₂₄F₃N₅O requir. 479.5 (free base). 25

Example 57

Procedure for the preparation of 3-methyl-2-methylthio-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone



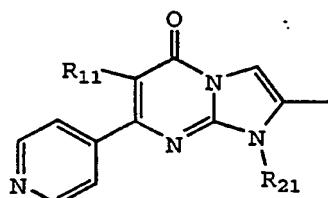
5 3-Methyl-2-methylthio-5-(3-methylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone: A solution of potassium t-butoxide (1M in t-butanol, 11, 1 mol) was added dropwise to a stirring solution of ethyl 3-methylphenyl acetate (178 g, 1 mol) in N,N-dimethylformamide (2 l). A solution of 10 4-cyanopyridine (104.11 g, 1 mol) in N,N-dimethylformamide (1 l) was pumped into the reaction mixture over a period of about 4.5 h. The mixture was then stirred at room temperature for 3 h, before the dropwise addition of a solution of methyl isothiocyanate 15 (68.4 ml, 1 mol) in N,N-dimethylformamide (50 ml) over a period of 10 min. After stirring for 1 h at room temperature, the reaction mixture was cooled to 3°C and methyl iodide (62.3 ml, 1 mol) was added dropwise over a period of 10 min. Stirring was continued at room 20 temperature overnight. The mixture was cooled to 3°C and water (4 l) was pumped into the reaction mixture over a period of 6 h. The precipitate was removed by filtration, washed with water and dried in a vacuum oven to give the title compound. MS (m/z): 324 (M+H)⁺; 25 C₁₈H₁₇N₃OS requir. 323.4.

Example 58

Using the corresponding starting materials, the following compounds of Table IV may be prepared using the procedure for 6-(4-fluorophenyl)-2-methyl-1-(3-30 phenylpropyl)-7-pyridin-4-yl-1*H*-imidazo(1,2-a)pyrimidin-5-one. The required pyrimidinones with the varied R"

substituents can be prepared using the general procedures described above.

TABLE IV



	R ₁₁	R ₂₁
5	3,5-dichlorophenyl 4-methoxyphenyl 3-tolyl 3-chlorophenyl	2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl
10	4-fluorophenyl 2-naphthyl n-butyl 2-thiophene 3-thiophene	2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl
15	3-aminophenyl 2-(5-chlorothiophene) 3-isopropylphenyl 3-tolyl 3-chlorophenyl	2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl
20	3-chloro-4-fluorophenyl 3,5-Ditrifluoromethylphenyl 4-fluorophenyl 3,4-dichlorophenyl 1-naphthyl	3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl
25	3-fluorophenyl 2-naphthyl n-butyl 2-thiophene 3-thiophene	3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl
30	3-aminophenyl 2-(5-chlorothiophene) 3,5-dichlorophenyl 4-tolyl 3-trifluoromethylphenyl	3-phenylpropyl 3-phenylpropyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl
35	4-methoxyphenyl 4-trifluoromethylphenyl 3-isopropylphenyl 3-tolyl 3-chlorophenyl	3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl
40	3-chloro-4-fluorophenyl 3,5-Ditrifluoromethylphenyl 4-fluorophenyl 3,4-dichlorophenyl	3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl

	2-naphthyl	3-methyl-3-phenyl-propyl
	n-butyl	3-methyl-3-phenyl-propyl
	2-thiophene	3-methyl-3-phenyl-propyl
	3-thiophene	3-methyl-3-phenyl-propyl
5	3-aminophenyl	3-methyl-3-phenyl-propyl
	2-(5-chlorothiophene)	3-methyl-3-phenyl-propyl
	3,5-dichlorophenyl	3-amino-3-phenyl-propyl
	4-tolyl	3-amino-3-phenyl-propyl
10	3-trifluoromethylphenyl	3-amino-3-phenyl-propyl
	4-methoxyphenyl	3-amino-3-phenyl-propyl
	4-trifluoromethylphenyl	3-amino-3-phenyl-propyl
	3-isopropylphenyl	3-amino-3-phenyl-propyl
	3-tolyl	3-amino-3-phenyl-propyl
	3-chlorophenyl	3-amino-3-phenyl-propyl
15	3-chloro-4-fluorophenyl	3-amino-3-phenyl-propyl
	3,5-Ditri fluoromethylphenyl	3-amino-3-phenyl-propyl
	4-fluorophenyl	3-amino-3-phenyl-propyl
	3,4-dichlorophenyl	3-amino-3-phenyl-propyl
	1-naphthyl	3-amino-3-phenyl-propyl
20	3-fluorophenyl	3-amino-3-phenyl-propyl
	2-naphthyl	3-amino-3-phenyl-propyl
	n-butyl	3-amino-3-phenyl-propyl
	2-thiophene	3-amino-3-phenyl-propyl
	3-thiophene	3-amino-3-phenyl-propyl
25	3-aminophenyl	3-amino-3-phenyl-propyl
	2-(5-chlorothiophene)	3-amino-3-phenyl-propyl
	3,5-dichlorophenyl	3-amino-3-phenyl-propyl
	4-tolyl	2(R)-amino-3-phenyl-propyl
30	3-trifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	4-methoxyphenyl	2(R)-amino-3-phenyl-propyl
	4-trifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	3-isopropylphenyl	2(R)-amino-3-phenyl-propyl
	3-tolyl	2(R)-amino-3-phenyl-propyl
	3-chlorophenyl	2(R)-amino-3-phenyl-propyl
35	3-chloro-4-fluorophenyl	2(R)-amino-3-phenyl-propyl
	3,5-Ditri fluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	4-fluorophenyl	2(R)-amino-3-phenyl-propyl
	3,4-dichlorophenyl	2(R)-amino-3-phenyl-propyl
	1-naphthyl	2(R)-amino-3-phenyl-propyl
40	3-fluorophenyl	2(R)-amino-3-phenyl-propyl
	2-naphthyl	2(R)-amino-3-phenyl-propyl
	n-butyl	2(R)-amino-3-phenyl-propyl
	2-thiophene	2(R)-amino-3-phenyl-propyl
	3-thiophene	2(R)-amino-3-phenyl-propyl
45	3-aminophenyl	2(R)-amino-3-phenyl-propyl
	2-(5-chlorothiophene)	2(R)-amino-3-phenyl-propyl
	3,5-dichlorophenyl	2-methyl-2-amino-3-phenyl-propyl
	4-tolyl	2-methyl-2-amino-3-phenyl-propyl
50	3-trifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
	4-methoxyphenyl	2-methyl-2-amino-3-phenyl-propyl
55	4-trifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl

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	3-isopropylphenyl	2-methyl-2-amino-3-phenyl-propyl
	3-tolyl	2-methyl-2-amino-3-phenyl-propyl
5	3-chlorophenyl	2-methyl-2-amino-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
10	3,5-Ditrifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
	4-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
	3,4-dichlorophenyl	2-methyl-2-amino-3-phenyl-propyl
15	1-naphthyl	2-methyl-2-amino-3-phenyl-propyl
	3-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
20	2-naphthyl	2-methyl-2-amino-3-phenyl-propyl
	n-butyl	2-methyl-2-amino-3-phenyl-propyl
	2-thiophene	2-methyl-2-amino-3-phenyl-propyl
25	3-thiophene	2-methyl-2-amino-3-phenyl-propyl
	3-aminophenyl	2-methyl-2-amino-3-phenyl-propyl
30	2-(5-chlorothiophene)	2-methyl-2-amino-3-phenyl-propyl
	3,5-dichlorophenyl	2-methyl-3-phenyl-propyl
	4-tolyl	2-methyl-3-phenyl-propyl
	3-trifluoromethylphenyl	2-methyl-3-phenyl-propyl
35	4-methoxyphenyl	2-methyl-3-phenyl-propyl
	4-trifluoromethylphenyl	2-methyl-3-phenyl-propyl
	3-isopropylphenyl	2-methyl-3-phenyl-propyl
	3-tolyl	2-methyl-3-phenyl-propyl
	3-chlorophenyl	2-methyl-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2-methyl-3-phenyl-propyl
40	3,5-Ditrifluoromethylphenyl	2-methyl-3-phenyl-propyl
	4-fluorophenyl	2-methyl-3-phenyl-propyl
	3,4-dichlorophenyl	2-methyl-3-phenyl-propyl
	1-naphthyl	2-methyl-3-phenyl-propyl
	3-fluorophenyl	2-methyl-3-phenyl-propyl
45	2-naphthyl	2-methyl-3-phenyl-propyl
	n-butyl	2-methyl-3-phenyl-propyl
	2-thiophene	2-methyl-3-phenyl-propyl
	3-thiophene	2-methyl-3-phenyl-propyl
	3-aminophenyl	2-methyl-3-phenyl-propyl
50	2-(5-chlorothiophene)	2-methyl-3-phenyl-propyl
	3,5-dichlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-tolyl	2-(N,N-dimethylamino)-3-phenyl-propyl
55	3-trifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl

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	4-methoxyphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-trifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
5	3-isopropylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-tolyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-chlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
10	3-chloro-4-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
15	4-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3,4-dichlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	1-naphthyl	2-(N,N-dimethylamino)-3-phenyl-propyl
20	3-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-naphthyl	2-(N,N-dimethylamino)-3-phenyl-propyl
25	n-butyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-thiophene	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-thiophene	2-(N,N-dimethylamino)-3-phenyl-propyl
30	3-aminophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-(5-chlorothiophene)	2-(N,N-dimethylamino)-3-phenyl-propyl
35	3,5-dichlorophenyl	2-(N-methylamino)-3-phenyl-propyl
	4-tolyl	2-(N-methylamino)-3-phenyl-propyl
	3-trifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
40	4-methoxyphenyl	2-(N-methylamino)-3-phenyl-propyl
	4-trifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
45	3-isopropylphenyl	2-(N-methylamino)-3-phenyl-propyl
	3-tolyl	2-(N-methylamino)-3-phenyl-propyl
	3-chlorophenyl	2-(N-methylamino)-3-phenyl-propyl
50	3-chloro-4-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
55	3,4-dichlorophenyl	2-(N-methylamino)-3-phenyl-propyl

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	4-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	1-naphthyl	2-(N-methylamino)-3-phenyl-propyl
5	3-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	2-naphthyl	2-(N-methylamino)-3-phenyl-propyl
10	n-butyl	2-(N-methylamino)-3-phenyl-propyl
	2-thiophene	2-(N-methylamino)-3-phenyl-propyl
	3-thiophene	2-(N-methylamino)-3-phenyl-propyl
15	3-aminophenyl	2-(N-methylamino)-3-phenyl-propyl
	2-(5-chlorothiophene)	2-(N-methylamino)-3-phenyl-propyl

Example 59

20 The compounds in table V can be prepared using the appropriate starting materials and the following procedures: The required pyrimidinones with the varied R¹¹ substituents can be prepared using the general procedures described above. The fused 6, 5 ring system can be prepared as described above affording R¹¹ as a hydrogen radical. Other R¹¹ groups can be introduced through a reductive amination process using the corresponding aldehyde with appropriate amino protection (Boc group). For example, N-Boc-phenylalanal can be prepared from the corresponding Weinreb amide through reduction with lithium aluminum hydride as described in the literature (Konieczny and Cushman Tetrahedron Lett 6939, 1992). The N-Boc-phenylalanal can then be reacted with the amino group using sodium triacetoxyborohydride.

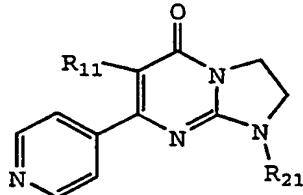
25 Alternatively, the alcohol of N-Boc-phenylalanol can be activated under Mitsunobu conditions (triphenylphosphine, diisopropyl azodicarboxylate) and reacted with the amino group of the 6, 5 fused system followed by removal of the Boc group (trifluoroacetic acid).

30

35

40

TABLE V



	R ₁₁	R ₂₁
5	3,5-dichlorophenyl 4-methoxyphenyl 3-tolyl 3-chlorophenyl 4-fluorophenyl 2-naphthyl	2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl
10	n-butyl 2-thiophene 3-thiophene 3-aminophenyl 2-(5-chlorothiophene)	2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl
15	3-isopropylphenyl 3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl 3,5-Ditri fluoromethylphenyl	3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl
20	4-fluorophenyl 3,4-dichlorophenyl 1-naphthyl 3-fluorophenyl 2-naphthyl	3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl
25	n-butyl 2-thiophene 3-thiophene 3-aminophenyl 2-(5-chlorothiophene)	3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl
30	3,5-dichlorophenyl 4-tolyl 3-trifluoromethylphenyl 4-methoxyphenyl 4-trifluoromethylphenyl	3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl
35	3-isopropylphenyl 3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl 3,5-Ditri fluoromethylphenyl	3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl
40	4-fluorophenyl 3,4-dichlorophenyl 2-naphthyl n-butyl 2-thiophene	3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl
45	3-thiophene 3-aminophenyl	3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl

	2-(5-chlorothiophene)	3-methyl-3-phenyl-propyl
	3,5-dichlorophenyl	3-amino-3-phenyl-propyl
	4-tolyl	3-amino-3-phenyl-propyl
	3-trifluoromethylphenyl	3-amino-3-phenyl-propyl
5	4-methoxyphenyl	3-amino-3-phenyl-propyl
	4-trifluoromethylphenyl	3-amino-3-phenyl-propyl
	3-isopropylphenyl	3-amino-3-phenyl-propyl
	3-tolyl	3-amino-3-phenyl-propyl
	3-chlorophenyl	3-amino-3-phenyl-propyl
10	3-chloro-4-fluorophenyl	3-amino-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	3-amino-3-phenyl-propyl
	4-fluorophenyl	3-amino-3-phenyl-propyl
	3,4-dichlorophenyl	3-amino-3-phenyl-propyl
	1-naphthyl	3-amino-3-phenyl-propyl
15	3-fluorophenyl	3-amino-3-phenyl-propyl
	2-naphthyl	3-amino-3-phenyl-propyl
	n-butyl	3-amino-3-phenyl-propyl
	2-thiophene	3-amino-3-phenyl-propyl
	3-thiophene	3-amino-3-phenyl-propyl
20	3-aminophenyl	3-amino-3-phenyl-propyl
	2-(5-chlorothiophene)	3-amino-3-phenyl-propyl
	3,5-dichlorophenyl	2(R)-amino-3-phenyl-propyl
	4-tolyl	2(R)-amino-3-phenyl-propyl
	3-trifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
25	4-methoxyphenyl	2(R)-amino-3-phenyl-propyl
	4-trifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	3-isopropylphenyl	2(R)-amino-3-phenyl-propyl
	3-tolyl	2(R)-amino-3-phenyl-propyl
	3-chlorophenyl	2(R)-amino-3-phenyl-propyl
30	3-chloro-4-fluorophenyl	2(R)-amino-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	4-fluorophenyl	2(R)-amino-3-phenyl-propyl
	3,4-dichlorophenyl	2(R)-amino-3-phenyl-propyl
	1-naphthyl	2(R)-amino-3-phenyl-propyl
35	3-fluorophenyl	2(R)-amino-3-phenyl-propyl
	2-naphthyl	2(R)-amino-3-phenyl-propyl
	n-butyl	2(R)-amino-3-phenyl-propyl
	2-thiophene	2(R)-amino-3-phenyl-propyl
	3-thiophene	2(R)-amino-3-phenyl-propyl
40	3-aminophenyl	2(R)-amino-3-phenyl-propyl
	2-(5-chlorothiophene)	2(R)-amino-3-phenyl-propyl
	3,5-dichlorophenyl	2(R)-amino-3-phenyl-propyl
	4-tolyl	2-methyl-2-amino-3-phenyl-propyl
45	3-trifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
	4-methoxyphenyl	2-methyl-2-amino-3-phenyl-propyl
50	4-trifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
	3-isopropylphenyl	2-methyl-2-amino-3-phenyl-propyl
	3-tolyl	2-methyl-2-amino-3-phenyl-propyl
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	3-chlorophenyl	2-methyl-2-amino-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
5	3,5-Ditrifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
	4-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
10	3,4-dichlorophenyl	2-methyl-2-amino-3-phenyl-propyl
	1-naphthyl	2-methyl-2-amino-3-phenyl-propyl
	3-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
15	2-naphthyl	2-methyl-2-amino-3-phenyl-propyl
	n-butyl	2-methyl-2-amino-3-phenyl-propyl
20	2-thiophene	2-methyl-2-amino-3-phenyl-propyl
	3-thiophene	2-methyl-2-amino-3-phenyl-propyl
	3-aminophenyl	2-methyl-2-amino-3-phenyl-propyl
25	2-(5-chlorothiophene)	2-methyl-2-amino-3-phenyl-propyl
	3,5-dichlorophenyl	2-methyl-3-phenyl-propyl
	4-tolyl	2-methyl-3-phenyl-propyl
30	3-trifluoromethylphenyl	2-methyl-3-phenyl-propyl
	4-methoxyphenyl	2-methyl-3-phenyl-propyl
	4-trifluoromethylphenyl	2-methyl-3-phenyl-propyl
	3-isopropylphenyl	2-methyl-3-phenyl-propyl
	3-tolyl	2-methyl-3-phenyl-propyl
	3-chlorophenyl	2-methyl-3-phenyl-propyl
35	3-chloro-4-fluorophenyl	2-methyl-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-methyl-3-phenyl-propyl
	4-fluorophenyl	2-methyl-3-phenyl-propyl
	3,4-dichlorophenyl	2-methyl-3-phenyl-propyl
40	1-naphthyl	2-methyl-3-phenyl-propyl
	3-fluorophenyl	2-methyl-3-phenyl-propyl
	2-naphthyl	2-methyl-3-phenyl-propyl
	n-butyl	2-methyl-3-phenyl-propyl
	2-thiophene	2-methyl-3-phenyl-propyl
45	3-thiophene	2-methyl-3-phenyl-propyl
	3-aminophenyl	2-methyl-3-phenyl-propyl
	2-(5-chlorothiophene)	2-methyl-3-phenyl-propyl
	3,5-dichlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-tolyl	2-(N,N-dimethylamino)-3-phenyl-propyl
50	3-trifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-methoxyphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
55	4-trifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl

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	3-isopropylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-tolyl	2-(N,N-dimethylamino)-3-phenyl-propyl
5	3-chlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
10	3,5-Ditrifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3,4-dichlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
15	1-naphthyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
20	2-naphthyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	n-butyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-thiophene	2-(N,N-dimethylamino)-3-phenyl-propyl
25	3-thiophene	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-aminophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-(5-chlorothiophene)	2-(N,N-dimethylamino)-3-phenyl-propyl
30	3,5-dichlorophenyl	2-(N-methylamino)-3-phenyl-propyl
	4-tolyl	2-(N-methylamino)-3-phenyl-propyl
35	3-trifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
	4-methoxyphenyl	2-(N-methylamino)-3-phenyl-propyl
	4-trifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
40	3-isopropylphenyl	2-(N-methylamino)-3-phenyl-propyl
	3-tolyl	2-(N-methylamino)-3-phenyl-propyl
45	3-chlorophenyl	2-(N-methylamino)-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
50	3,4-dichlorophenyl	2-(N-methylamino)-3-phenyl-propyl
	4-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
55	1-naphthyl	2-(N-methylamino)-3-phenyl-propyl

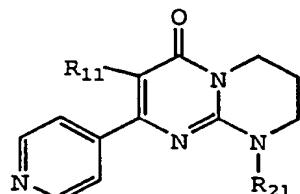
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	3-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	2-naphthyl	2-(N-methylamino)-3-phenyl-propyl
5	n-butyl	2-(N-methylamino)-3-phenyl-propyl
	2-thiophene	2-(N-methylamino)-3-phenyl-propyl
10	3-thiophene	2-(N-methylamino)-3-phenyl-propyl
	3-aminophenyl	2-(N-methylamino)-3-phenyl-propyl
	2-(5-chlorothiophene)	2-(N-methylamino)-3-phenyl-propyl

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Example 60

The compounds in table VI can be prepared using the appropriate starting materials and procedures as described above.

TABLE VI

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	R ₁₁	R ₂₁
	3,5-dichlorophenyl	2(S)-amino-3-phenyl-propyl
	4-methoxyphenyl	2(S)-amino-3-phenyl-propyl
	3-tolyl	2(S)-amino-3-phenyl-propyl
25	3-chlorophenyl	2(S)-amino-3-phenyl-propyl
	4-fluorophenyl	2(S)-amino-3-phenyl-propyl
	2-naphthyl	2(S)-amino-3-phenyl-propyl
	n-butyl	2(S)-amino-3-phenyl-propyl
	2-thiophene	2(S)-amino-3-phenyl-propyl
30	3-thiophene	2(S)-amino-3-phenyl-propyl
	3-aminophenyl	2(S)-amino-3-phenyl-propyl
	2-(5-chlorothiophene)	2(S)-amino-3-phenyl-propyl
	3-isopropylphenyl	3-phenylpropyl
	3-tolyl	3-phenylpropyl
35	3-chlorophenyl	3-phenylpropyl
	3-chloro-4-fluorophenyl	3-phenylpropyl
	3,5-Difluoromethylphenyl	3-phenylpropyl
	4-fluorophenyl	3-phenylpropyl
	3,4-dichlorophenyl	3-phenylpropyl
40	1-naphthyl	3-phenylpropyl
	3-fluorophenyl	3-phenylpropyl
	2-naphthyl	3-phenylpropyl
	n-butyl	3-phenylpropyl

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	2-thiophene	3-phenylpropyl
	3-thiophene	3-phenylpropyl
	3-aminophenyl	3-phenylpropyl
	2-(5-chlorothiophene)	3-phenylpropyl
5	3,5-dichlorophenyl	3-methyl-3-phenyl-propyl
	4-tolyl	3-methyl-3-phenyl-propyl
	3-trifluoromethylphenyl	3-methyl-3-phenyl-propyl
	4-methoxyphenyl	3-methyl-3-phenyl-propyl
	4-trifluoromethylphenyl	3-methyl-3-phenyl-propyl
10	3-isopropylphenyl	3-methyl-3-phenyl-propyl
	3-tolyl	3-methyl-3-phenyl-propyl
	3-chlorophenyl	3-methyl-3-phenyl-propyl
	3-chloro-4-fluorophenyl	3-methyl-3-phenyl-propyl
	3,5-Ditri fluoromethylphenyl	3-methyl-3-phenyl-propyl
15	4-fluorophenyl	3-methyl-3-phenyl-propyl
	3,4-dichlorophenyl	3-methyl-3-phenyl-propyl
	2-naphthyl	3-methyl-3-phenyl-propyl
	n-butyl	3-methyl-3-phenyl-propyl
	2-thiophene	3-methyl-3-phenyl-propyl
20	3-thiophene	3-methyl-3-phenyl-propyl
	3-aminophenyl	3-methyl-3-phenyl-propyl
	2-(5-chlorothiophene)	3-methyl-3-phenyl-propyl
	3,5-dichlorophenyl	3-amino-3-phenyl-propyl
	4-tolyl	3-amino-3-phenyl-propyl
25	3-trifluoromethylphenyl	3-amino-3-phenyl-propyl
	4-methoxyphenyl	3-amino-3-phenyl-propyl
	4-trifluoromethylphenyl	3-amino-3-phenyl-propyl
	3-isopropylphenyl	3-amino-3-phenyl-propyl
	3-tolyl	3-amino-3-phenyl-propyl
30	3-chlorophenyl	3-amino-3-phenyl-propyl
	3-chloro-4-fluorophenyl	3-amino-3-phenyl-propyl
	3,5-Ditri fluoromethylphenyl	3-amino-3-phenyl-propyl
	4-fluorophenyl	3-amino-3-phenyl-propyl
	3,4-dichlorophenyl	3-amino-3-phenyl-propyl
35	1-naphthyl	3-amino-3-phenyl-propyl
	3-fluorophenyl	3-amino-3-phenyl-propyl
	2-naphthyl	3-amino-3-phenyl-propyl
	n-butyl	3-amino-3-phenyl-propyl
	2-thiophene	3-amino-3-phenyl-propyl
40	3-thiophene	3-amino-3-phenyl-propyl
	3-aminophenyl	3-amino-3-phenyl-propyl
	2-(5-chlorothiophene)	3-amino-3-phenyl-propyl
	3,5-dichlorophenyl	2(R)-amino-3-phenyl-propyl
	4-tolyl	2(R)-amino-3-phenyl-propyl
45	3-trifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	4-methoxyphenyl	2(R)-amino-3-phenyl-propyl
	4-trifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	3-isopropylphenyl	2(R)-amino-3-phenyl-propyl
	3-tolyl	2(R)-amino-3-phenyl-propyl
50	3-chlorophenyl	2(R)-amino-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2(R)-amino-3-phenyl-propyl
	3,5-Ditri fluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	4-fluorophenyl	2(R)-amino-3-phenyl-propyl
	3,4-dichlorophenyl	2(R)-amino-3-phenyl-propyl
55	1-naphthyl	2(R)-amino-3-phenyl-propyl
	3-fluorophenyl	2(R)-amino-3-phenyl-propyl

	2-naphthyl	2 (R)-amino-3-phenyl-propyl
	n-butyl	2 (R)-amino-3-phenyl-propyl
	2-thiophene	2 (R)-amino-3-phenyl-propyl
	3-thiophene	2 (R)-amino-3-phenyl-propyl
5	3-aminophenyl	2 (R)-amino-3-phenyl-propyl
	2-(5-chlorothiophene)	2 (R)-amino-3-phenyl-propyl
	3,5-dichlorophenyl	2-methyl-2-amino-3-phenyl-propyl
	4-tolyl	2-methyl-2-amino-3-phenyl-propyl
10	3-trifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
	4-methoxyphenyl	2-methyl-2-amino-3-phenyl-propyl
15	4-trifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
	3-isopropylphenyl	2-methyl-2-amino-3-phenyl-propyl
	3-tolyl	2-methyl-2-amino-3-phenyl-propyl
20	3-chlorophenyl	2-methyl-2-amino-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
25	3,5-Ditrifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
	4-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
	3,4-dichlorophenyl	2-methyl-2-amino-3-phenyl-propyl
30	1-naphthyl	2-methyl-2-amino-3-phenyl-propyl
	3-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
35	2-naphthyl	2-methyl-2-amino-3-phenyl-propyl
	n-butyl	2-methyl-2-amino-3-phenyl-propyl
	2-thiophene	2-methyl-2-amino-3-phenyl-propyl
40	3-thiophene	2-methyl-2-amino-3-phenyl-propyl
	3-aminophenyl	2-methyl-2-amino-3-phenyl-propyl
45	2-(5-chlorothiophene)	2-methyl-2-amino-3-phenyl-propyl
	3,5-dichlorophenyl	2-methyl-3-phenyl-propyl
	4-tolyl	2-methyl-3-phenyl-propyl
	3-trifluoromethylphenyl	2-methyl-3-phenyl-propyl
50	4-methoxyphenyl	2-methyl-3-phenyl-propyl
	4-trifluoromethylphenyl	2-methyl-3-phenyl-propyl
	3-isopropylphenyl	2-methyl-3-phenyl-propyl
	3-tolyl	2-methyl-3-phenyl-propyl
	3-chlorophenyl	2-methyl-3-phenyl-propyl
55	3-chloro-4-fluorophenyl	2-methyl-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-methyl-3-phenyl-propyl

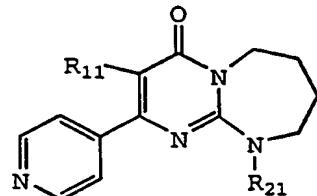
	4-fluorophenyl	2-methyl-3-phenyl-propyl
	3,4-dichlorophenyl	2-methyl-3-phenyl-propyl
	1-naphthyl	2-methyl-3-phenyl-propyl
	3-fluorophenyl	2-methyl-3-phenyl-propyl
5	2-naphthyl	2-methyl-3-phenyl-propyl
	n-butyl	2-methyl-3-phenyl-propyl
	2-thiophene	2-methyl-3-phenyl-propyl
	3-thiophene	2-methyl-3-phenyl-propyl
	3-aminophenyl	2-methyl-3-phenyl-propyl
10	2-(5-chlorothiophene)	2-methyl-3-phenyl-propyl
	3,5-dichlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-tolyl	2-(N,N-dimethylamino)-3-phenyl-propyl
15	3-trifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-methoxyphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-trifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
20	3-isopropylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-tolyl	2-(N,N-dimethylamino)-3-phenyl-propyl
25	3-chlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
30	4-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3,4-dichlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
35	1-naphthyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-naphthyl	2-(N,N-dimethylamino)-3-phenyl-propyl
40	n-butyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-thiophene	2-(N,N-dimethylamino)-3-phenyl-propyl
45	3-thiophene	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-aminophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-(5-chlorothiophene)	2-(N,N-dimethylamino)-3-phenyl-propyl
50	3,5-dichlorophenyl	2-(N-methylamino)-3-phenyl-propyl
	4-tolyl	2-(N-methylamino)-3-phenyl-propyl
55	3-trifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl

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	4-methoxyphenyl	2-(N-methylamino)-3-phenyl-propyl
	4-trifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
5	3-isopropylphenyl	2-(N-methylamino)-3-phenyl-propyl
	3-tolyl	2-(N-methylamino)-3-phenyl-propyl
	3-chlorophenyl	2-(N-methylamino)-3-phenyl-propyl
10	3-chloro-4-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
15	3,4-dichlorophenyl	2-(N-methylamino)-3-phenyl-propyl
	4-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	1-naphthyl	2-(N-methylamino)-3-phenyl-propyl
20	3-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	2-naphthyl	2-(N-methylamino)-3-phenyl-propyl
25	n-butyl	2-(N-methylamino)-3-phenyl-propyl
	2-thiophene	2-(N-methylamino)-3-phenyl-propyl
	3-thiophene	2-(N-methylamino)-3-phenyl-propyl
30	3-aminophenyl	2-(N-methylamino)-3-phenyl-propyl
	2-(5-chlorothiophene)	2-(N-methylamino)-3-phenyl-propyl

35 **Example 61**

The compounds in table VII can be prepared using the appropriate starting materials and procedures as described above.

TABLE VII

40

R ₁₁	R ₂₁
3,5-dichlorophenyl	2(S)-amino-3-phenyl-propyl
4-methoxyphenyl	2(S)-amino-3-phenyl-propyl
3-tolyl	2(S)-amino-3-phenyl-propyl

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	3-chlorophenyl	2(S)-amino-3-phenyl-propyl
	4-fluorophenyl	2(S)-amino-3-phenyl-propyl
	2-naphthyl	2(S)-amino-3-phenyl-propyl
	n-butyl	2(S)-amino-3-phenyl-propyl
5	2-thiophene	2(S)-amino-3-phenyl-propyl
	3-thiophene	2(S)-amino-3-phenyl-propyl
	3-aminophenyl	2(S)-amino-3-phenyl-propyl
	2-(5-chlorothiophene)	2(S)-amino-3-phenyl-propyl
	3-isopropylphenyl	2(S)-amino-3-phenyl-propyl
10	3-tolyl	3-phenylpropyl
	3-chlorophenyl	3-phenylpropyl
	3-chloro-4-fluorophenyl	3-phenylpropyl
	3,5-Ditrifluoromethylphenyl	3-phenylpropyl
	4-fluorophenyl	3-phenylpropyl
15	3,4-dichlorophenyl	3-phenylpropyl
	1-naphthyl	3-phenylpropyl
	3-fluorophenyl	3-phenylpropyl
	2-naphthyl	3-phenylpropyl
	n-butyl	3-phenylpropyl
20	2-thiophene	3-phenylpropyl
	3-thiophene	3-phenylpropyl
	3-aminophenyl	3-phenylpropyl
	2-(5-chlorothiophene)	3-phenylpropyl
	3,5-dichlorophenyl	3-methyl-3-phenyl-propyl
25	4-tolyl	3-methyl-3-phenyl-propyl
	3-trifluoromethylphenyl	3-methyl-3-phenyl-propyl
	4-methoxyphenyl	3-methyl-3-phenyl-propyl
	4-trifluoromethylphenyl	3-methyl-3-phenyl-propyl
	3-isopropylphenyl	3-methyl-3-phenyl-propyl
30	3-tolyl	3-methyl-3-phenyl-propyl
	3-chlorophenyl	3-methyl-3-phenyl-propyl
	3-chloro-4-fluorophenyl	3-methyl-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	3-methyl-3-phenyl-propyl
	4-fluorophenyl	3-methyl-3-phenyl-propyl
35	3,4-dichlorophenyl	3-methyl-3-phenyl-propyl
	2-naphthyl	3-methyl-3-phenyl-propyl
	n-butyl	3-methyl-3-phenyl-propyl
	2-thiophene	3-methyl-3-phenyl-propyl
	3-thiophene	3-methyl-3-phenyl-propyl
40	3-aminophenyl	3-methyl-3-phenyl-propyl
	2-(5-chlorothiophene)	3-methyl-3-phenyl-propyl
	3,5-dichlorophenyl	3-amino-3-phenyl-propyl
	4-tolyl	3-amino-3-phenyl-propyl
	3-trifluoromethylphenyl	3-amino-3-phenyl-propyl
45	4-methoxyphenyl	3-amino-3-phenyl-propyl
	4-trifluoromethylphenyl	3-amino-3-phenyl-propyl
	3-isopropylphenyl	3-amino-3-phenyl-propyl
	3-tolyl	3-amino-3-phenyl-propyl
	3-chlorophenyl	3-amino-3-phenyl-propyl
50	3-chloro-4-fluorophenyl	3-amino-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	3-amino-3-phenyl-propyl
	4-fluorophenyl	3-amino-3-phenyl-propyl
	3,4-dichlorophenyl	3-amino-3-phenyl-propyl
	1-naphthyl	3-amino-3-phenyl-propyl
55	3-fluorophenyl	3-amino-3-phenyl-propyl
	2-naphthyl	3-amino-3-phenyl-propyl

	n-butyl	3-amino-3-phenyl-propyl
	2-thiophene	3-amino-3-phenyl-propyl
	3-thiophene	3-amino-3-phenyl-propyl
	3-aminophenyl	3-amino-3-phenyl-propyl
5	2-(5-chlorothiophene)	3-amino-3-phenyl-propyl
	3,5-dichlorophenyl	2(R)-amino-3-phenyl-propyl
	4-tolyl	2(R)-amino-3-phenyl-propyl
	3-trifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	4-methoxyphenyl	2(R)-amino-3-phenyl-propyl
10	4-trifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	3-isopropylphenyl	2(R)-amino-3-phenyl-propyl
	3-tolyl	2(R)-amino-3-phenyl-propyl
	3-chlorophenyl	2(R)-amino-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2(R)-amino-3-phenyl-propyl
15	3,5-Ditrifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	4-fluorophenyl	2(R)-amino-3-phenyl-propyl
	3,4-dichlorophenyl	2(R)-amino-3-phenyl-propyl
	1-naphthyl	2(R)-amino-3-phenyl-propyl
	3-fluorophenyl	2(R)-amino-3-phenyl-propyl
20	2-naphthyl	2(R)-amino-3-phenyl-propyl
	n-butyl	2(R)-amino-3-phenyl-propyl
	2-thiophene	2(R)-amino-3-phenyl-propyl
	3-thiophene	2(R)-amino-3-phenyl-propyl
	3-aminophenyl	2(R)-amino-3-phenyl-propyl
25	2-(5-chlorothiophene)	2(R)-amino-3-phenyl-propyl
	3,5-dichlorophenyl	2-methyl-2-amino-3-phenyl-propyl
	4-tolyl	2-methyl-2-amino-3-phenyl-propyl
30	3-trifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
	4-methoxyphenyl	2-methyl-2-amino-3-phenyl-propyl
	4-trifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
35	3-isopropylphenyl	2-methyl-2-amino-3-phenyl-propyl
	3-tolyl	2-methyl-2-amino-3-phenyl-propyl
40	3-chlorophenyl	2-methyl-2-amino-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
45	4-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
	3,4-dichlorophenyl	2-methyl-2-amino-3-phenyl-propyl
50	1-naphthyl	2-methyl-2-amino-3-phenyl-propyl
	3-fluorophenyl	2-methyl-2-amino-3-phenyl-propyl
55	2-naphthyl	2-methyl-2-amino-3-phenyl-propyl

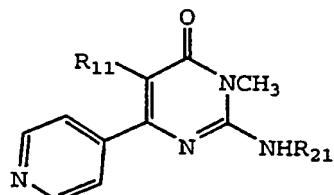
	n-butyl	2-methyl-2-amino-3-phenyl-propyl
	2-thiophene	2-methyl-2-amino-3-phenyl-propyl
5	3-thiophene	2-methyl-2-amino-3-phenyl-propyl
	3-aminophenyl	2-methyl-2-amino-3-phenyl-propyl
	2-(5-chlorothiophene)	2-methyl-2-amino-3-phenyl-propyl
10	3,5-dichlorophenyl	2-methyl-3-phenyl-propyl
	4-tolyl	2-methyl-3-phenyl-propyl
	3-trifluoromethylphenyl	2-methyl-3-phenyl-propyl
	4-methoxyphenyl	2-methyl-3-phenyl-propyl
15	4-trifluoromethylphenyl	2-methyl-3-phenyl-propyl
	3-isopropylphenyl	2-methyl-3-phenyl-propyl
	3-tolyl	2-methyl-3-phenyl-propyl
	3-chlorophenyl	2-methyl-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2-methyl-3-phenyl-propyl
20	3,5-Ditrifluoromethylphenyl	2-methyl-3-phenyl-propyl
	4-fluorophenyl	2-methyl-3-phenyl-propyl
	3,4-dichlorophenyl	2-methyl-3-phenyl-propyl
	1-naphthyl	2-methyl-3-phenyl-propyl
	3-fluorophenyl	2-methyl-3-phenyl-propyl
25	2-naphthyl	2-methyl-3-phenyl-propyl
	n-butyl	2-methyl-3-phenyl-propyl
	2-thiophene	2-methyl-3-phenyl-propyl
	3-thiophene	2-methyl-3-phenyl-propyl
	3-aminophenyl	2-methyl-3-phenyl-propyl
30	2-(5-chlorothiophene)	2-methyl-3-phenyl-propyl
	3,5-dichlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-tolyl	2-(N,N-dimethylamino)-3-phenyl-propyl
35	3-trifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-methoxyphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-trifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
40	3-isopropylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-tolyl	2-(N,N-dimethylamino)-3-phenyl-propyl
45	3-chlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
50	4-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3,4-dichlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
55	1-naphthyl	2-(N,N-dimethylamino)-3-phenyl-propyl

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	3-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-naphthyl	2-(N,N-dimethylamino)-3-phenyl-propyl
5	n-butyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-thiophene	2-(N,N-dimethylamino)-3-phenyl-propyl
10	3-thiophene	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-aminophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-(5-chlorothiophene)	2-(N,N-dimethylamino)-3-phenyl-propyl
15	3,5-dichlorophenyl	2-(N-methylamino)-3-phenyl-propyl
	4-tolyl	2-(N-methylamino)-3-phenyl-propyl
20	3-trifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
	4-methoxyphenyl	2-(N-methylamino)-3-phenyl-propyl
	4-trifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
25	3-isopropylphenyl	2-(N-methylamino)-3-phenyl-propyl
	3-tolyl	2-(N-methylamino)-3-phenyl-propyl
	3-chlorophenyl	2-(N-methylamino)-3-phenyl-propyl
30	3-chloro-4-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
35	3,4-dichlorophenyl	2-(N-methylamino)-3-phenyl-propyl
	4-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
40	1-naphthyl	2-(N-methylamino)-3-phenyl-propyl
	3-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	2-naphthyl	2-(N-methylamino)-3-phenyl-propyl
45	n-butyl	2-(N-methylamino)-3-phenyl-propyl
	2-thiophene	2-(N-methylamino)-3-phenyl-propyl
	3-thiophene	2-(N-methylamino)-3-phenyl-propyl
50	3-aminophenyl	2-(N-methylamino)-3-phenyl-propyl
	2-(5-chlorothiophene)	2-(N-methylamino)-3-phenyl-propyl

Example 62

Using the corresponding starting materials, the following compounds of Table VIII may be prepared using the procedure for 3-methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone.

TABLE VIII

	R_{11}	R_{21}
10	3,5-dichlorophenyl 4-methoxyphenyl 3-tolyl 3-chlorophenyl 4-fluorophenyl	2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl
15	2-naphthyl n-butyl 2-thiophene 3-thiophene 3-aminophenyl	2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl 2(S)-amino-3-phenyl-propyl
20	2-(5-chlorothiophene) 3-isopropylphenyl 3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl	2(S)-amino-3-phenyl-propyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl
25	3,5-Ditrifluoromethylphenyl 4-fluorophenyl 3,4-dichlorophenyl 1-naphthyl 3-fluorophenyl	3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl
30	2-naphthyl n-butyl 2-thiophene 3-thiophene 3-aminophenyl	3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl 3-phenylpropyl
35	2-(5-chlorothiophene) 3,5-dichlorophenyl 4-tolyl 3-trifluoromethylphenyl 4-methoxyphenyl	3-phenylpropyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl
40	4-trifluoromethylphenyl 3-isopropylphenyl 3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl	3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl 3-methyl-3-phenyl-propyl

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	3,5-Ditrifluoromethylphenyl	3-methyl-3-phenyl-propyl
	4-fluorophenyl	3-methyl-3-phenyl-propyl
	3,4-dichlorophenyl	3-methyl-3-phenyl-propyl
	2-naphthyl	3-methyl-3-phenyl-propyl
5	n-butyl	3-methyl-3-phenyl-propyl
	2-thiophene	3-methyl-3-phenyl-propyl
	3-thiophene	3-methyl-3-phenyl-propyl
	3-aminophenyl	3-methyl-3-phenyl-propyl
	2-(5-chlorothiophene)	3-methyl-3-phenyl-propyl
10	3,5-dichlorophenyl	3-amino-3-phenyl-propyl
	4-tolyl	3-amino-3-phenyl-propyl
	3-trifluoromethylphenyl	3-amino-3-phenyl-propyl
	4-methoxyphenyl	3-amino-3-phenyl-propyl
	4-trifluoromethylphenyl	3-amino-3-phenyl-propyl
15	3-isopropylphenyl	3-amino-3-phenyl-propyl
	3-tolyl	3-amino-3-phenyl-propyl
	3-chlorophenyl	3-amino-3-phenyl-propyl
	3-chloro-4-fluorophenyl	3-amino-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	3-amino-3-phenyl-propyl
20	4-fluorophenyl	3-amino-3-phenyl-propyl
	3,4-dichlorophenyl	3-amino-3-phenyl-propyl
	1-naphthyl	3-amino-3-phenyl-propyl
	3-fluorophenyl	3-amino-3-phenyl-propyl
	2-naphthyl	3-amino-3-phenyl-propyl
25	n-butyl	3-amino-3-phenyl-propyl
	2-thiophene	3-amino-3-phenyl-propyl
	3-thiophene	3-amino-3-phenyl-propyl
	3-aminophenyl	3-amino-3-phenyl-propyl
	2-(5-chlorothiophene)	3-amino-3-phenyl-propyl
30	3,5-dichlorophenyl	2(R)-amino-3-phenyl-propyl
	4-tolyl	2(R)-amino-3-phenyl-propyl
	3-trifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
	4-methoxyphenyl	2(R)-amino-3-phenyl-propyl
	4-trifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
35	3-isopropylphenyl	2(R)-amino-3-phenyl-propyl
	3-tolyl	2(R)-amino-3-phenyl-propyl
	3-chlorophenyl	2(R)-amino-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2(R)-amino-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2(R)-amino-3-phenyl-propyl
40	4-fluorophenyl	2(R)-amino-3-phenyl-propyl
	3,4-dichlorophenyl	2(R)-amino-3-phenyl-propyl
	1-naphthyl	2(R)-amino-3-phenyl-propyl
	3-fluorophenyl	2(R)-amino-3-phenyl-propyl
	2-naphthyl	2(R)-amino-3-phenyl-propyl
45	n-butyl	2(R)-amino-3-phenyl-propyl
	2-thiophene	2(R)-amino-3-phenyl-propyl
	3-thiophene	2(R)-amino-3-phenyl-propyl
	3-aminophenyl	2(R)-amino-3-phenyl-propyl
	2-(5-chlorothiophene)	2(R)-amino-3-phenyl-propyl
50	3,5-dichlorophenyl	2-methyl-2-amino-3-phenyl-propyl
	4-tolyl	2-methyl-2-amino-3-phenyl-propyl
	3-trifluoromethylphenyl	2-methyl-2-amino-3-phenyl-propyl
55		

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	4-methoxyphenyl	2-methyl-2-amino-3-phenyl-
	4-trifluoromethylphenyl	propyl
5	3-isopropylphenyl	2-methyl-2-amino-3-phenyl-
	3-tolyl	propyl
	3-chlorophenyl	2-methyl-2-amino-3-phenyl-
10	3-chloro-4-fluorophenyl	propyl
	3,5-Ditrifluoromethylphenyl	2-methyl-2-amino-3-phenyl-
15	4-fluorophenyl	propyl
	3,4-dichlorophenyl	2-methyl-2-amino-3-phenyl-
	1-naphthyl	propyl
20	3-fluorophenyl	2-methyl-2-amino-3-phenyl-
	2-naphthyl	propyl
25	n-butyl	2-methyl-2-amino-3-phenyl-
	2-thiophene	propyl
	3-thiophene	2-methyl-2-amino-3-phenyl-
30	3-aminophenyl	propyl
	2-(5-chlorothiophene)	2-methyl-2-amino-3-phenyl-
35	3,5-dichlorophenyl	propyl
	4-tolyl	2-methyl-3-phenyl-propyl
	3-trifluoromethylphenyl	2-methyl-3-phenyl-propyl
	4-methoxyphenyl	2-methyl-3-phenyl-propyl
	4-trifluoromethylphenyl	2-methyl-3-phenyl-propyl
40	3-isopropylphenyl	2-methyl-3-phenyl-propyl
	3-tolyl	2-methyl-3-phenyl-propyl
	3-chlorophenyl	2-methyl-3-phenyl-propyl
	3-chloro-4-fluorophenyl	2-methyl-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-methyl-3-phenyl-propyl
45	4-fluorophenyl	2-methyl-3-phenyl-propyl
	3,4-dichlorophenyl	2-methyl-3-phenyl-propyl
	1-naphthyl	2-methyl-3-phenyl-propyl
	3-fluorophenyl	2-methyl-3-phenyl-propyl
	2-naphthyl	2-methyl-3-phenyl-propyl
50	n-butyl	2-methyl-3-phenyl-propyl
	2-thiophene	2-methyl-3-phenyl-propyl
	3-thiophene	2-methyl-3-phenyl-propyl
	3-aminophenyl	2-methyl-3-phenyl-propyl
	2-(5-chlorothiophene)	2-methyl-3-phenyl-propyl
55	3,5-dichlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl

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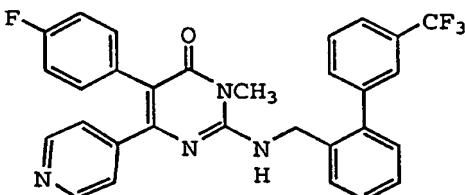
	4-tolyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-trifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
5	4-methoxyphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	4-trifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
10	3-isopropylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-tolyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-chlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
15	3-chloro-4-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3,5-Ditrifluoromethylphenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
20	4-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	3,4-dichlorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	1-naphthyl	2-(N,N-dimethylamino)-3-phenyl-propyl
25	3-fluorophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-naphthyl	2-(N,N-dimethylamino)-3-phenyl-propyl
30	n-butyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-thiophene	2-(N,N-dimethylamino)-3-phenyl-propyl
	3-thiophene	2-(N,N-dimethylamino)-3-phenyl-propyl
35	3-aminophenyl	2-(N,N-dimethylamino)-3-phenyl-propyl
	2-(5-chlorothiophene)	2-(N,N-dimethylamino)-3-phenyl-propyl
40	3,5-dichlorophenyl	2-(N-methylamino)-3-phenyl-propyl
	4-tolyl	2-(N-methylamino)-3-phenyl-propyl
	3-trifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
45	4-methoxyphenyl	2-(N-methylamino)-3-phenyl-propyl
	4-trifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
50	3-isopropylphenyl	2-(N-methylamino)-3-phenyl-propyl
	3-tolyl	2-(N-methylamino)-3-phenyl-propyl
	3-chlorophenyl	2-(N-methylamino)-3-phenyl-propyl
55	3-chloro-4-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl

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	3,5-Ditrifluoromethylphenyl	2-(N-methylamino)-3-phenyl-propyl
	3,4-dichlorophenyl	2-(N-methylamino)-3-phenyl-propyl
5	4-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	1-naphthyl	2-(N-methylamino)-3-phenyl-propyl
10	3-fluorophenyl	2-(N-methylamino)-3-phenyl-propyl
	2-naphthyl	2-(N-methylamino)-3-phenyl-propyl
	n-butyl	2-(N-methylamino)-3-phenyl-propyl
15	2-thiophene	2-(N-methylamino)-3-phenyl-propyl
	3-thiophene	2-(N-methylamino)-3-phenyl-propyl
	3-aminophenyl	2-(N-methylamino)-3-phenyl-propyl
20	2-(5-chlorothiophene)	2-(N-methylamino)-3-phenyl-propyl

Example 63

Procedure for the preparation of 2-((2-(3-trifluoromethylphenyl)phenylmethyl)amino)-3-methyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone



Step A. 2-((2-bromophenylmethyl)amino)-5-(4-fluorophenyl)-6-(4-pyridyl)-3-methyl-4(3H)-pyrimidinone:

30 The compound, 3-methyl-5-(4-fluorophenyl)-6-(4-pyridyl)-2-thiomethyl-4(3H)-pyrimidinone (470 mg, 1.44 mmol) was dissolved in methanol:water mixture(1.8:1, 40ml and 22.5ml). Potassium peroxymonosulfate (OXONE Aldrich Chem Co., 2.5g 4.1 mmol) was added to a cooled (4°C) reaction mixture and then the reaction was continued for 16h at room-temperature. The reaction mixture was concentrated and extracted with dichloromethane and the organic layer was washed with water, dried over Na₂SO₄, and was concentrated. The residue (500mg) and o-

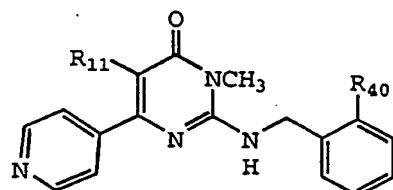
35

bromobenzylamine were mixed in 1,4-dioxane (20 ml). The clear solution was heated at 85°C for 18 h and progress of the reaction monitored by TLC. The reaction mixture was concentrated and chromatographed on a silica gel column to obtain the titled compound. MS(m/z): 466.9 C₂₃H₁₈BrFN₄O requirs: 465.33 1H-NMR (CDCl₃): d 8.49 (dd, 2H, pyridyl), 7.67-6.81 (m, 12H, Ph and pyridyl), 5.44 (t, 1H, NH), 4.92 (d 2H, CH₂-Ph), 3.6 (s, 3H, N-CH₃).

Step B. 2-((2-(3-trifluoromethylphenyl)phenylmethyl)amino)-3-methyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone: 2-((2-bromophenylmethyl)amino)-5-(4-fluorophenyl)-6-(4-pyridyl)-3-methyl-4(3H)-pyrimidinone (175 mg, 0.38 mmol) was dispersed in 2M sodium carbonate solution (12 ml) and 3-trifluoromethylbenzene- boronic acid (170 mg, 0.89 mmol), toluene (12ml) were added to the above mixture and the reaction mixture was degassed and catalyst tetrakis(triphenylphosphine) Pd(0) (50 mg) was added. The reaction mixture was refluxed for 16 h. The formation of the product was monitored by TLC. Then it was cooled, diluted with toluene (12 ml) and washed with water. The organic layer was dried over sodium sulfate, concentrated and the product was purified by silica gel chromatography to give the title compound. MS(m/z): 531.1 C₂₃H₂₂F₄N₄O requir. 530.53; 1H-NMR (CDCl₃): d 8.43 (m, 2H, pyridyl), 7.69-7.12 (m, 8H, Ph), 7.11-6.88 (m, 6H, pyridyl and Ph-CF₃), 4.85 (m, 3H, CH₂-Ph and NH), 3.32 (N-CH₃).

Example 64

Using the corresponding starting materials, the following compounds of Table IX may be prepared using the procedure for 2-((2-(3-trifluoromethylphenyl)phenylmethyl)amino)-3-methyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone.

TABLE IX

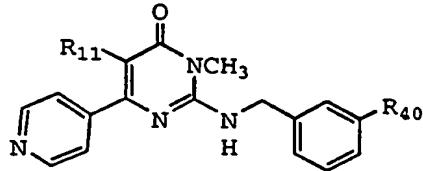
	R_{11}	R_{40}
5	4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl	3,5-dichlorophenyl 4-tolyl 4-methoxyphenyl 4-trifluoromethylphenyl 3-isopropylphenyl 3-tolyl 3-chlorophenyl
10	4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl	3-chloro-4-fluorophenyl 3,5-Ditrifluoromethylphenyl 4-fluorophenyl 3,4-dichlorophenyl
15	4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl	1-naphthyl 3-fluorophenyl 2-naphthyl n-butyl 2-thiophene
20	4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl	3-thiophene 3-aminophenyl 2-(5-chlorothiophene) 3,5-dichlorophenyl 4-tolyl 3-trifluoromethylphenyl
25	3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl	4-methoxyphenyl 4-trifluoromethylphenyl 3-isopropylphenyl 3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl
30	3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl	3,5-Ditrifluoromethylphenyl 4-fluorophenyl 3,4-dichlorophenyl
35	3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl	1-naphthyl 3-fluorophenyl 2-naphthyl n-butyl 2-thiophene
40	3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl	3-thiophene 3-aminophenyl 2-(5-chlorothiophene)

Example 65

Using the corresponding starting materials, the
45 following compounds of Table X may be prepared using the

procedure for 2-((2-(3-trifluoromethylphenyl)phenylmethyl)amino)-3-methyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone.

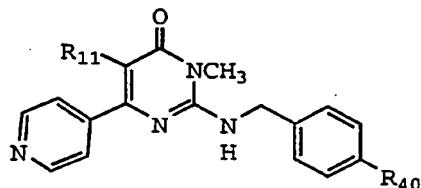
TABLE X



	R_{11}	R_{40}
5		
10	4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl	3,5-dichlorophenyl 4-tolyl 4-methoxyphenyl 4-trifluoromethylphenyl 3-isopropylphenyl 3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl 3,5-Ditrifluoromethylphenyl
15	4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl	4-fluorophenyl 3,4-dichlorophenyl 1-naphthyl 3-fluorophenyl 2-naphthyl n-butyl 2-thiophene 3-thiophene 3-aminophenyl
20	4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl	2-(5-chlorothiophene) 3,5-dichlorophenyl 4-tolyl 3-trifluoromethylphenyl 4-methoxyphenyl 4-trifluoromethylphenyl 3-isopropylphenyl 3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl 3,5-Ditrifluoromethylphenyl
25	4-fluorophenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl	4-fluorophenyl 3,4-dichlorophenyl 1-naphthyl 3-fluorophenyl 2-naphthyl n-butyl 2-thiophene 3-thiophene 3-aminophenyl
30	3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl	3,5-Ditrifluoromethylphenyl 4-trifluoromethylphenyl 3-isopropylphenyl 3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl 3,5-Ditrifluoromethylphenyl 4-fluorophenyl 3,4-dichlorophenyl 1-naphthyl 3-fluorophenyl 2-naphthyl n-butyl 2-thiophene 3-thiophene 3-aminophenyl
35	3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl	2-(5-chlorothiophene) 3,5-dichlorophenyl 4-tolyl 3-trifluoromethylphenyl 4-methoxyphenyl 4-trifluoromethylphenyl 3-isopropylphenyl 3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl 3,5-Ditrifluoromethylphenyl 4-fluorophenyl 3,4-dichlorophenyl 1-naphthyl 3-fluorophenyl 2-naphthyl n-butyl 2-thiophene 3-thiophene 3-aminophenyl
40	3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl	2-(5-chlorothiophene) 3,5-dichlorophenyl 4-tolyl 3-trifluoromethylphenyl 4-methoxyphenyl 4-trifluoromethylphenyl 3-isopropylphenyl 3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl 3,5-Ditrifluoromethylphenyl 4-fluorophenyl 3,4-dichlorophenyl 1-naphthyl 3-fluorophenyl 2-naphthyl n-butyl 2-thiophene 3-thiophene 3-aminophenyl
45	3-trifluoromethylphenyl	2-(5-chlorothiophene)

Example 66

Using the corresponding starting materials, the following compounds of Table XI may be prepared using the procedure for 2-((2-(3-trifluoromethylphenyl)phenylmethyl)amino)-3-methyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone.

TABLE XI

	R_{11}	R_{40}
10	4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl	3,5-dichlorophenyl 4-tolyl 4-methoxyphenyl 4-trifluoromethylphenyl 3-isopropylphenyl
15	4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl	3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl 3,5-Ditri fluoromethylphenyl 4-fluorophenyl
20	4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl	3,4-dichlorophenyl 1-naphthyl 3-fluorophenyl 2-naphthyl n-butyl
25	4-fluorophenyl 4-fluorophenyl 4-fluorophenyl 4-fluorophenyl	2-thiophene 3-thiophene 3-aminophenyl 2-(5-chlorothiophene)
30	3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl	3,5-dichlorophenyl 4-tolyl 3-trifluoromethylphenyl 4-methoxyphenyl 4-trifluoromethylphenyl
35	3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl	3-isopropylphenyl 3-tolyl 3-chlorophenyl 3-chloro-4-fluorophenyl 3,5-Ditri fluoromethylphenyl
40	3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl 3-trifluoromethylphenyl	4-fluorophenyl 3,4-dichlorophenyl 1-naphthyl 3-fluorophenyl 2-naphthyl n-butyl
45	3-trifluoromethylphenyl	2-thiophene

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3-trifluoromethylphenyl	3-thiophene
3-trifluoromethylphenyl	3-aminophenyl
3-trifluoromethylphenyl	2-(5-chlorothiophene)

Example 67

5

Biological Assays

The following assays were used to characterize the ability of compounds of the invention to inhibit the production of TNF- α and IL-1- β . The second assay measured the inhibition of TNF- α and/or IL-1- β in mice after oral administration of the test compounds. The third assay, a glucagon binding inhibition *in vitro* assay, can be used to characterize the ability of compounds of the invention to inhibit glucagon binding. The fourth assay, a Cyclooxygenase enzyme (COX-1 and COX-2) inhibition activity *in vitro* assay, can be used to characterize the ability of compounds of the invention to inhibit COX-1 and/or COX-2.

Lipopolysaccharide-activated monocyte TNF production assay20 **Isolation of monocytes**

Test compounds were evaluated *in vitro* for the ability to inhibit the production of TNF by monocytes activated with bacterial lipopolysaccharide (LPS). Fresh residual source leukocytes (a byproduct of plateletpheresis) were obtained from a local blood bank, and peripheral blood mononuclear cells (PBMCs) were isolated by density gradient centrifugation on Ficol-Paque Plus (Pharmacia). PBMCs were suspended at 2 x 10⁶/ml in DMEM supplemented to contain 2% FCS, 10 mM, 0.3 mg/ml glutamate, 100 U/ml penicillin G and 100 mg/ml streptomycin sulfate (complete media). Cells were plated into Falcon flat bottom, 96 well culture plates (200 μ l/well) and cultured overnight at 37°C and 6% CO₂. Non-adherent cells were removed by washing with 200 μ l/well of fresh medium. Wells containing adherent cells (~70% monocytes) were replenished with 100 μ l of fresh medium.

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Preparation of test compound stock solutions

Test compounds were dissolved in DMZ. Compound stock solutions were prepared to an initial concentration of 10 - 50 μ M. Stocks were diluted initially to 20 - 200 μ M in complete media. Nine two-fold serial dilutions of each compound were then prepared in complete medium.

10 One hundred microliters of each test compound dilution were added to microtiter wells containing adherent monocytes and 100 μ l complete medium. Monocytes were cultured with test compounds for 60 min at which time 25 μ l of complete medium containing 30 ng/ml lipopolysaccharide from *E. coli* K532 were added to each well. Cells were cultured an additional 4 hrs. Culture supernatants were then removed and TNF presence in the supernatants was quantified using an ELISA.

15 *TNF ELISA*
20 Flat bottom, 96 well Corning High Binding ELISA plates were coated overnight (4°C) with 150 μ L/well of 3 μ g/ml murine anti-human TNF- α MAb (R&D Systems #MAB210). Wells were then blocked for 1 hr at room temperature with 200 μ L/well of CaCl₂-free ELISA buffer supplemented 25 to contain 20 mg/ml BSA (standard ELISA buffer: 20 mM, 150 mM NaCl, 2 mM CaCl₂, 0.15 mM thimerosal, pH 7.4). Plates were washed and replenished with 100 μ l of test supernatants (diluted 1:3) or standards. Standards consisted of eleven 1.5-fold serial dilutions from a 30 stock of 1 ng/ml recombinant human TNF (R&D Systems). Plates were incubated at room temperature for 1 hr on orbital shaker (300 rpm), washed and replenished with 100 μ l/well of 0.5 μ g/ml goat anti-human TNF- α (R&D systems #AB-210-NA) biotinylated at a 4:1 ratio. Plates 35 were incubated for 40 min, washed and replenished with 100 μ l/well of alkaline phosphatase-conjugated

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streptavidin (Jackson ImmunoResearch #016-050-084) at 0.02 µg/ml. Plates were incubated 30 min, washed and replenished with 200 µl/well of 1 mg/ml of p-nitrophenyl phosphate. After 30 min, plates were read at 405 nm on 5 a V_{max} plate reader.

Data analysis

Standard curve data were fit to a second order polynomial and unknown TNF-α concentrations determined from their OD by solving this equation for 10 concentration. TNF concentrations were then plotted vs. test compound concentration using a second order polynomial. This equation was then used to calculate the concentration of test compounds causing a 50% reduction in TNF production.

15 Compounds of the invention can also be shown to inhibit LPS-induced release of IL-1β, IL-6 and/or IL-8 from monocytes by measuring concentrations of IL-1β, IL-6 and/or IL-8 by methods well known to those skilled in the art. In a similar manner to the above described 20 assay involving the LPS induced release of TNF-α from monocytes, compounds of this invention can also be shown to inhibit LPS induced release of IL-1β, IL-6 and/or IL-8 from monocytes by measuring concentrations of IL-1β, IL-6 and/or IL-8 by methods well known to those skilled 25 in the art. Thus, the compounds of the invention may lower elevated levels of TNF-α, IL-1, IL-6, and IL-8 levels. Reducing elevated levels of these inflammatory cytokines to basal levels or below is favorable in controlling, slowing progression, and alleviating many 30 disease states. All of the compounds are useful in the methods of treating disease states in which TNF-α, IL-1β, IL-6, and IL-8 play a role to the full extent of the definition of TNF-α-mediated diseases described herein.

Inhibition of LPS-Induced TNF-α production in mice

Male DBA/1LACJ mice were dosed with vehicle or test compounds in a vehicle (the vehicle consisting of 0.5% tragacanth in 0.03 N HCl) 30 minutes prior to lipopolysaccharide (2 mg/kg, I.V.) injection. Ninety 5 minutes after LPS injection, blood was collected and the serum was analyzed by ELISA for TNF levels.

The following compounds exhibit activities in the monocyte assay (LPS induced TNF release) with IC₅₀ values of 20 μM or less:

10 2-(2,6-Dichlorobenzyl)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
2-(Butylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
15 2-(Benzylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
5-(4-Fluorophenyl)-3-methyl-((R-1-phenylethyl)amino)-(4-pyridyl)-4(3H)-pyrimidinone
20 2-(2-Chlorophenyl)-ethylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
5-(4-Fluorophenyl)-2-(2-(4-fluorophenyl)-ethylamino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
5-(4-Fluorophenyl)-2-((2-hydroxy-2-phenyl)-ethylamino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
25 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
5-(4-Fluorophenyl)-3-methyl-2-((1-methyl-3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
5-(4-Fluorophenyl)-3-methyl-2-((R-1-methyl-3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
30 5-(4-Fluorophenyl)-3-methyl-2-((2-phenylaminoethyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
5-(4-Fluorophenyl)-2-((3-imidazolylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
35 5-(4-Fluorophenyl)-3-methyl-6-(4-pyridyl)-2-(3-pyrrolidin-1-yl)-propylamino)-4(3H)-pyrimidinone
3,6-Diphenyl-4-(4-pyridyl)-2(1H)-pyridone
6-(4-Methylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone
40 6-(4-Ethylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone
6-(2,4-Dimethylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone
3-Phenyl-4-(4-pyridyl)-6-(2-thienyl)-2(1H)-pyridone
6-(2-Furyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone

2-(((S)-2-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 2-(((R)-2-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 5 2-((S)-2-N-Ethyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 2-((2-Amino-2-methyl-3-phenylpropyl)amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 10 2-((2-Aminomethyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 2-((3-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 15 5-(4-Fluorophenyl)-3-methyl-2-(3-(2-methylphenyl)propyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
 5-(4-Fluorophenyl)-3-methyl-2-((R,S)-2-amino-3-(2'-fluorophenyl)-propyl-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
 20 2-((S)-2-Acetamido-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 5-(4-Fluorophenyl)-2-((S)-2-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 25 2-((S)-2-N-n-Butylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 2-((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 5-(4-Fluorophenyl)-3-methyl-2-((2-methyl-3-phenylpropyl)amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
 30 2-((S)-2-Amino-3-phenylpropyl)-amino)-3-ethyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone
 3-Ethyl-5-(4-fluorophenyl)-2-((2-methyl-3-phenylpropyl)amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
 35 2-((2-(3-trifluoromethylphenyl)phenylmethyl)amino)-3-methyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone
 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(4-tolyl)-6-(4-pyridyl)-4(3H)-pyrimidinone
 40 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(4-trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone
 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-isopropylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone
 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-chloro-4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3,5-bis(trifluoromethyl)phenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

5 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3,4-dichlorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(1-naphthyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

10 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(3-phenylpropylamino)-5-(3,5-dichlorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

15 3-Methyl-2-(3-phenylpropylamino)-5-(4-tolyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(3-phenylpropylamino)-5-(3-trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(3-phenylpropylamino)-5-(4-methoxyphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

20 3-Methyl-2-(3-phenylpropylamino)-5-(4-trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(2-methyl-3-phenylpropylamino)-5-(3-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

25 3-Methyl-2-(2-methyl-3-phenylpropylamino)-5-(1-naphthyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

5-(4-Fluorophenyl)-2-(((S)-2-N-glycylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

30 2-(((S)-2-N-Glycylamino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone

5-(4-Fluorophenyl)-2-(((S)-2-hydroxyacetamido-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

35 5-(4-Fluorophenyl)-2-(((S)-2-pyrrolidinyl-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

2-((S)-3-Benzylpiperazinyl)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

40 2-((3-Amino-3-(2-fluorophenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

2-((3-Amino-3-(2-methylphenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

2-(((S)-3-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

45 2-(((R)-3-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

2-(((S)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone

2-(((R)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone

5 2-((3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone

2-((3-Amino-3-(2-methylphenyl)propyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone

10 2-((3-Amino-3-(2-fluorophenyl)propyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone

2-((3-Amino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone

15 2-((3-Amino-3-(2-fluorophenyl)propyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone

2-((3-Amino-3-(2-chlorophenyl)propyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone

20 2-(((S)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3,4-dimethylphenyl)-4-(3H)-pyrimidinone

2-(((2R,3R)-3-Amino-2-methyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

25 2-(((2S,3S)-3-Amino-2-methyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

5-(4-Fluorophenyl)-2-(((S)-3-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

30 5-(4-Fluorophenyl)-2-(((R)-3-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone

5-(4-Fluorophenyl)-3-methyl-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-4-(3H)-pyrimidinone

35 3-Methyl-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone

3-Methyl-5-(3-methylphenyl)-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-4-(3H)-pyrimidinone

40 3-Methyl-5-(4-methylthiophenyl)-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-4-(3H)-pyrimidinone

45 2-(((S)-2-Amino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone

5-(4-Fluorophenyl)-2-((3-hydroxy-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 2-(((S)-2-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
 5 2-(((S)-2-Amino-3-(2-fluorophenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 2-(((S)-2-Amino-3-(4-fluorophenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 10 2-(((S)-2-Amino-3-(2-chlorophenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 2-(((S)-2-N-Isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone
 15 2-(((S)-2-N-Isopropylamino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
 5-(3-Chlorophenyl)-2-(((S)-2-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 20 2-(((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
 2-(((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-3-methyl-5-(3-chlorophenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
 25 2-(((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-5-(3-trifluorophenyl)-4-(3H)-pyrimidinone
 5-(4-Fluorophenyl)-3-methyl-2-(((S)-2-N-methylamino-3-phenylpropyl)-amino)-6-(4-pyridyl)-4-(3H)-pyrimidinone.
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The following compounds exhibit activities in the monocyte assay (LPS induced TNF release) with IC₅₀ values of 5 μM or less:

35 2-(2,6-Dichlorobenzyl)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 2-(Benzylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 40 5-(4-Fluorophenyl)-3-methyl-((R-1-phenylethyl)amino)-(4-pyridyl)-4(3H)-pyrimidinone
 2-(2-(2-Chlorophenyl)-ethylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 5-(4-Fluorophenyl)-2-(2-(4-fluorophenyl)-ethylamino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 45 5-(4-Fluorophenyl)-3-methyl-2-(3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone

5-(4-Fluorophenyl)-3-methyl-2-((1-methyl-3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
 5-(4-Fluorophenyl)-3-methyl-2-((R-1-methyl-3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
 5 5-(4-Fluorophenyl)-3-methyl-2-((2-phenylaminoethyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
 5-(4-Fluorophenyl)-3-methyl-6-(4-pyridyl)-2-(3-(pyrrolidin-1-yl)-propylamino)-4(3H)-pyrimidinone
 6-(4-Ethylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone
 10 2-((S)-2-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 2-((R)-2-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 15 2-((S)-2-N-Ethyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 2-((2-Amino-2-methoxy-3-phenylpropyl) amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 2-((2-Aminomethyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 20 2-((3-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 5-(4-Fluorophenyl)-3-methyl-2-(3-(2-methylphenyl)propyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
 25 5-(4-Fluorophenyl)-3-methyl-2-((R,S)-2-amino-3-(2'-fluorophenyl)-propyl-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
 2-((S)-2-Acetamido-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 30 5-(4-Fluorophenyl)-2-((S)-2-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 2-((S)-2-N-n-Butylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 35 2-((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone
 5-(4-Fluorophenyl)-3-methyl-2-((2-methoxy-3-phenylpropyl) amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
 40 2-((S)-2-Amino-3-phenylpropyl)-amino)-3-ethyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone
 3-Ethyl-5-(4-fluorophenyl)-2-((2-methoxy-3-phenylpropyl) amino)-6-(4-pyridyl)-4(3H)-pyrimidinone
 2-((2-(3-trifluoromethylphenyl)phenylmethyl)amino)-3-methyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone
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3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(4-tolyl)-
6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(4-
trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

5 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-
isopropylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-chloro-
4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

10 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3,5-
bis(trifluoromethyl)phenyl)-6-(4-pyridyl)-4(3H)-
pyrimidinone

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3,4-
dichlorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

15 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(1-
naphthyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-
fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-
trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

20 3-Methyl-2-(3-phenylpropylamino)-5-(3,5-dichlorophenyl)-
6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(3-phenylpropylamino)-5-(4-tolyl)-6-(4-
pyridyl)-4(3H)-pyrimidinone

25 3-Methyl-2-(3-phenylpropylamino)-5-(3-
trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(3-phenylpropylamino)-5-(4-methoxyphenyl)-6-
(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(3-phenylpropylamino)-5-(4-
trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

30 3-Methyl-2-(2-methyl-3-phenylpropylamino)-5-(3-
fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

3-Methyl-2-(2-methyl-3-phenylpropylamino)-5-(1-
naphthyl)-6-(4-pyridyl)-4(3H)-pyrimidinone

35 5-(4-Fluorophenyl)-2-(((S)-2-N-glycylamino-3-
phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-
pyrimidinone

2-(((S)-2-N-Glycylamino-3-phenylpropyl)-amino)-3-methyl-
5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone

40 5-(4-Fluorophenyl)-2-(((S)-2-hydroxyacetamido-3-
phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-
pyrimidinone

5-(4-Fluorophenyl)-2-(((S)-2-pyrrolidinyl-3-
phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-
pyrimidinone

45 2-((S)-3-Benzylpiperazinyl)-5-(4-fluorophenyl)-3-methyl-
6-(4-pyridyl)-4-(3H)-pyrimidinone

2-((3-Amino-3-(2-fluorophenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 2-((3-Amino-3-(2-methylphenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 5 2-((S)-3-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 2-((R)-3-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 10 2-((S)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone
 2-((R)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone
 2-((3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone
 15 2-((3-Amino-3-(2-methylphenyl)propyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone
 2-((3-Amino-3-(2-fluorophenyl)propyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone
 20 2-((3-Amino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
 2-((3-Amino-3-(2-fluorophenyl)propyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
 25 2-((3-Amino-3-(2-chlorophenyl)propyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
 2-((S)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3,4-dimethylphenyl)-4-(3H)-pyrimidinone
 30 2-(((2R,3R)-3-Amino-2-methyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 2-(((2S,3S)-3-Amino-2-methyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 35 5-(4-Fluorophenyl)-2-((S)-3-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 5-(4-Fluorophenyl)-2-((R)-3-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
 40 5-(4-Fluorophenyl)-3-methyl-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-4-(3H)-pyrimidinone
 3-Methyl-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone
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3-Methyl-5-(3-methylphenyl)-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-4-(3H)-pyrimidinone
5 3-Methyl-5-(4-methylthiophenyl)-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-4-(3H)-pyrimidinone
10 2-((S)-2-Amino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
15 5-(4-Fluorophenyl)-2-((3-hydroxy-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
20 2-((S)-2-Amino-3-(2-fluorophenyl)propyl)-amino)-5-(4-fluorophenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
25 2-((S)-2-Amino-3-(2-chlorophenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
30 2-((S)-2-N-Isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone
35 2-((S)-2-N-Isopropylamino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
40 5-(3-Chlorophenyl-2-((S)-2-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone
45 2-((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
2-((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-3-methyl-5-(3-chlorophenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone
2-((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-5-(3-trifluorophenyl)-4-(3H)-pyrimidinone
5-(4-Fluorophenyl)-3-methyl-2-((S)-2-N-methylamino-3-phenylpropyl)-amino)-6-(4-pyridyl)-4-(3H)-pyrimidinone
40 Compounds of the invention may be shown to have anti-inflammatory properties in animal models of inflammation, including carageenan paw edema, collagen induced arthritis and adjuvant arthritis, such as the carageenan paw edema model (C. A. Winter et al Proc. Soc. Exp. Biol. Med. (1962) vol 111, p 544; K. F.

Swingle, in R. A. Scherrer and M. W. Whitehouse, Eds.,
Antiinflammatory Agents, Chemistry and Pharmacology,
Vol. 13-II, Academic, New York, 1974, p. 33) and
collagen induced arthritis (D. E. Trentham et al J. Exp.
5 Med. (1977) vol. 146, p 857; J. S. Courtenay, Nature
(New Biol.) (1980), Vol 283, p 666).

¹²⁵I-Glucagon Binding Screen with CHO/hGLUR Cells

The assay is described in WO 97/16442, which is
10 incorporated herein by reference in its entirety.

Reagents

The reagents can be prepared as follows: (a) prepare fresh 1M o-Phenanthroline (Aldrich) (198.2 mg/ml ethanol); (b) prepare fresh 0.5M DTT (Sigma); (c) 15 Protease Inhibitor Mix (1000X): 5 mg leupeptin, 10 mg benzamidine, 40 mg bacitracin and 5 mg soybean trypsin inhibitor per ml DMSO and store aliquots at -20°C; (d) 250 µM human glucagon (Peninsula): solubilize 0.5 mg vial in 575 µl 0.1N acetic acid (1 µl yields 1 µM final concentration in assay for non-specific binding) and store in aliquots at -20°C; (e) Assay Buffer: 20mM Tris (pH 7.8), 1 mM DTT and 3 mM o-phenanthroline; (f) Assay Buffer with 0.1% BSA (for dilution of label only; 0.01% final in assay): 10 µl 10% BSA (heat-inactivated) and 25 990 µl Assay Buffer; (g) ¹²⁵I-Glucagon (NEN, receptor-grade, 2200 Ci/mmol): dilute to 50,000 cpm/25 µl in assay buffer with BSA (about 50pM final concentration in assay).

Harvesting of CHO/hGLUR Cells for Assay

30 1. Remove media from confluent flask then rinse once each with PBS (Ca, Mg-free) and Enzyme-free Dissociation Fluid (Specialty Media, Inc.).

2. Add 10 ml Enzyme-free Dissoc. Fluid and hold for about 4 min. at 37°C.

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3. Gently tap cells free, triturate, take aliquot for counting and centrifuge remainder for 5 min. at 1000 rpm.

4. Resuspend pellet in Assay Buffer at 75000 cells per 100 µl.

Membrane preparations of CHO/hGLUR cells can be used in place of whole cells at the same assay volume. Final protein concentration of a membrane preparation is determined on a per batch basis.

10 Assay

The determination of inhibition of glucagon binding can be carried out by measuring the reduction of I^{125} -glucagon binding in the presence of compounds of Formula I. The reagents are combined as follows:

15

	Compound/ Vehicle	250 µM Glucagon	I^{125} - Glucagon	CHO/hGLUR Cells
Total Binding	--/5 µl	--	25 µl	100 µl
+ Compound	5 µl/--	--	25 µl	100 µl
Nonspecific Binding	--/5 µl	1 µl	25 µl	100 µl

The mixture is incubated for 60 min. at 22°C on a shaker at 275 rpm. The mixture is filtered over pre-soaked (0.5% polyethylinime (PEI)) GF/C filtermat using an 20 Innotech Harvester or Tomtec Harvester with four washes of ice-cold 20mM Tris buffer (pH 7.8). The radioactivity in the filters is determined by a gamma-scintillation counter.

Thus, compounds of the invention may also be shown 25 to inhibit the binding of glucagon to glucagon receptors.

Cyclooxygenase Enzyme Activity Assay

The human monocytic leukemia cell line, THP-1, differentiated by exposure to phorbol esters expresses only COX-1; the human osteosarcoma cell line 143B expresses predominantly COX-2. THP-1 cells are 5 routinely cultured in RPMI complete media supplemented with 10% FBS and human osteosarcoma cells (HOSC) are cultured in minimal essential media supplemented with 10% fetal bovine serum (MEM-10%FBS); all cell incubations are at 37°C in a humidified environment 10 containing 5% CO₂.

COX-1 Assay

In preparation for the COX-1 assay, THP-1 cells are grown to confluence, split 1:3 into RPMI containing 2% 15 FBS and 10 mM phorbol 12-myristate 13-acetate (TPA), and incubated for 48 hours on a shaker to prevent attachment. Cells are pelleted and resuspended in Hank's Buffered Saline (HBS) at a concentration of 2.5 × 20 10⁶ cells/mL and plated in 96-well culture plates at a density of 5 × 10³ cells/mL. Test compounds are diluted in HBS and added to the desired final concentration and the cells are incubated for an additional 4 hours. Arachidonic acid is added to a final concentration of 30 mM, the cells incubated for 20 minutes at 37°C, and 25 enzyme activity determined as described below.

COX-2 Assay

For the COX-2 assay, subconfluent HOSC are trypsinized and resuspended at 3 × 10⁶ cells/mL in MEM- 30 FBS containing 1 ng human IL-1b/mL, plated in 96-well tissue culture plates at a density of 3 × 10⁴ cells per well, incubated on a shaker for 1 hour to evenly distribute cells, followed by an additional 2 hour static incubation to allow attachment. The media is 35 then replaced with MEM containing 2% FBS (MEM-2%FBS) and 1 ng human IL-1b/mL, and the cells incubated for 18-22

hours. Following replacement of media with 190 mL MEM, 10 mL of test compound diluted in HBS is added to achieve the desired concentration and the cells incubated for 4 hours. The supernatants are removed and 5 replaced with MEM containing 30 mM arachidonic acid, the cells incubated for 20 minutes at 37°C, and enzyme activity determined as described below.

COX Activity Determined

10 After incubation with arachidonic acid, the reactions are stopped by the addition of 1 N HCl, followed by neutralization with 1 N NaOH and centrifugation to pellet cell debris. Cyclooxygenase enzyme activity in both HOSC and THP-1 cell supernatants 15 is determined by measuring the concentration of PGE, using a commercially available ELISA (Neogen #404110). A standard curve of PGE, is used for calibration, and commercially available COX-1 and COX-2 inhibitors are included as standard controls.

20 Accordingly, the compounds of the invention or a pharmaceutical composition thereof are useful for prophylaxis and treatment of rheumatoid arthritis; Pagets disease; osteophorosis; multiple myeloma; uveitis; acute and chronic myelogenous leukemia; 25 pancreatic β cell destruction; osteoarthritis; rheumatoid spondylitis; gouty arthritis; inflammatory bowel disease; adult respiratory distress syndrome (ARDS); psoriasis; Crohn's disease; allergic rhinitis; ulcerative colitis; anaphylaxis; contact dermatitis; 30 asthma; muscle degeneration; cachexia; Reiter's syndrome; type I and type II diabetes; bone resorption diseases; graft vs. host reaction; ischemia reperfusion injury; atherosclerosis; brain trauma; Alzheimer's disease; stroke; myocardial infarction; multiple 35 sclerosis; cerebral malaria; sepsis; septic shock; toxic shock syndrome; fever, and myalgias due to infection. HIV-1, HIV-2, HIV-3, cytomegalovirus (CMV), influenza,

adenovirus, the herpes viruses (including HSV-1, HSV-2), and herpes zoster, all of which are sensitive to TNF- α and/or IL-1 inhibition or glucagon antagonism, will also be positively effected by the compounds and methods of
5 the invention.

The compounds of the present invention also may possess analgesic properties and may be useful for the treatment of pain disorders, such as hyperalgesia due to excessive IL-1. The compounds of the present invention
10 may also prevent the production of prostaglandins by inhibition of enzymes in the human arachidonic acid/prostaglandin pathway, including cyclooxygenase (WO 96/03387, incorporated herein by reference in its entirety).

15 Because of their ability to lower TNF- α and IL-1 concentrations or inhibit glucagon binding to its receptor, the compounds of the invention are also useful research tools for studying the physiology associated with blocking these effects.

20 The methods of the invention comprise administering an effective dose of a compound of the invention, a pharmaceutical salt thereof, or a pharmaceutical composition of either, to a subject (i.e., an animal, preferably a mammal, most preferably a human) in need of
25 a reduction in the level of TNF- α , IL-1, IL-6, and/or IL-8 levels and/or reduction in plasma glucose levels and/or which subject may be suffering from rheumatoid arthritis; Pagets disease; osteoporosis; multiple myeloma; uveitis; acute and chronic myelogenous
30 leukemia; pancreatic & cell destruction; osteoarthritis; rheumatoid spondylitis; gouty arthritis; inflammatory bowel disease; adult respiratory distress syndrome (ARDS); psoriasis; Crohn's disease; allergic rhinitis; ulcerative colitis; anaphylaxis; contact dermatitis;
35 asthma; muscle degeneration; cachexia; Reiter's syndrome; type I and type II diabetes; bone resorption diseases; graft vs. host reaction; Alzheimer's disease;

stroke; myocardial infarction; ischemia reperfusion injury; atherosclerosis; brain trauma; multiple sclerosis; cerebral malaria; sepsis; septic shock; toxic shock syndrome; fever, and myalgias due to infection, or 5 which subject is infected by HIV-1, HIV-2, HIV-3, cytomegalovirus (CMV), influenza, adenovirus, the herpes viruses (including HSV-1, HSV-2), or herpes zoster.

In another aspect, this invention comprises the use of a compound of the invention, or pharmaceutically 10 acceptable salts thereof, in the manufacture of a medicament for the treatment either acutely or chronically of a TNF- α , IL-1 β , IL-6, and/or IL-8 mediated disease state, including those described previously. Also, the compounds of this invention are useful in the 15 manufacture of a analgesic medicament and a medicament for treating pain disorders, such as hyperalgesia. The compounds of the present invention also are useful in the manufacture of a medicament to prevent the production of prostaglandins by inhibition of enzymes in the human 20 arachidonic acid/prostaglandin pathway.

In still another aspect, this invention provides a pharmaceutical composition comprising an effective TNF- α , IL-1 β , IL-6, and/or IL-8 lowering amount and/or effective plasma glucose level lowering amount of a 25 compound of the invention and a pharmaceutically acceptable carrier or diluent, and if desired other active ingredients. The compounds of the invention are administered by any suitable route, preferably in the form of a pharmaceutical composition adapted to such a 30 route, and in a dose effective for the treatment intended. Therapeutically effective doses of the compounds of the present invention required to arrest the progress or prevent tissue damage associated with the disease are readily ascertained by one of ordinary 35 skill in the art using standard methods.

For the treatment of TNF- α , IL-1 β , IL-6, and IL-8 mediated diseases and/or hyperglycemia, the compounds of

the present invention may be administered orally, parentally, by inhalation spray, rectally, or topically in dosage unit formulations containing conventional pharmaceutically acceptable carriers, adjuvants, and 5 vehicles. The term parenteral as used herein includes, subcutaneous, intravenous, intramuscular, intrasternal, infusion techniques or intraperitoneally.

The dosage regimen for treating a TNF- α , IL-1, IL-6, and IL-8 mediated diseases and/or hyperglycemia with 10 the compounds of this invention and/or compositions of this invention is based on a variety of factors, including the type of disease, the age, weight, sex, medical condition of the patient, the severity of the condition, the route of administration, and the 15 particular compound employed. Thus, the dosage regimen may vary widely, but can be determined routinely using standard methods. Dosage levels of the order from about 0.01 mg to 30 mg per kilogram of body weight per day, preferably from about 0.1 mg to 10 mg/kg, more 20 preferably from about 0.25 mg to 1 mg/kg are useful for all methods of use disclosed herein.

The pharmaceutically active compounds of this invention can be processed in accordance with conventional methods of pharmacy to produce medicinal 25 agents for administration to patients, including humans and other mammals.

For oral administration, the pharmaceutical composition may be in the form of, for example, a capsule, a tablet, a suspension, or liquid. The 30 pharmaceutical composition is preferably made in the form of a dosage unit containing a given amount of the active ingredient. For example, these may contain an amount of active ingredient from about 1 to 2000 mg, preferably from about 1 to 500 mg, more preferably from 35 about 5 to 150 mg. A suitable daily dose for a human or other mammal may vary widely depending on the condition

of the patient and other factors, but, once again, can be determined using routine methods.

The active ingredient may also be administered by injection as a composition with suitable carriers
5 including saline, dextrose, or water. The daily parenteral dosage regimen will be from about 0.1 to about 30 mg/kg of total body weight, preferably from about 0.1 to about 10 mg/kg, and more preferably from about 0.25 mg to 1 mg/kg.

10 Injectable preparations, such as sterile injectable aqueous or oleaginous suspensions, may be formulated according to the known art using suitable dispersing or wetting agents and suspending agents. The sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally acceptable diluent or solvent, for example as a solution in 1,3-butanediol. Among the acceptable vehicles and solvents that may be employed are water, Ringer's solution, and isotonic sodium chloride solution. In
15 addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose any bland fixed oil may be employed, including synthetic mono- or diglycerides. In addition, fatty acids such as oleic acid find use in the preparation of
20 injectables.
25

Suppositories for rectal administration of the drug can be prepared by mixing the drug with a suitable non-irritating excipient such as cocoa butter and polyethylene glycols that are solid at ordinary
30 temperatures but liquid at the rectal temperature and will therefore melt in the rectum and release the drug.

A suitable topical dose of active ingredient of a compound of the invention is 0.1 mg to 150 mg administered one to four, preferably one or two times
35 daily. For topical administration, the active ingredient may comprise from 0.001% to 10% w/w, e.g., from 1% to 2% by weight of the formulation, although it

may comprise as much as 10% w/w, but preferably not more than 5% w/w, and more preferably from 0.1% to 1% of the formulation.

Formulations suitable for topical administration
5 include liquid or semi-liquid preparations suitable for penetration through the skin (e.g., liniments, lotions, ointments, creams, or pastes) and drops suitable for administration to the eye, ear, or nose.

For administration, the compounds of this invention
10 are ordinarily combined with one or more adjuvants appropriate for the indicated route of administration. The compounds may be admixed with lactose, sucrose, starch powder, cellulose esters of alcanoic acids, stearic acid, talc, magnesium stearate, magnesium oxide,
15 sodium and calcium salts of phosphoric and sulphuric acids, acacia, gelatin, sodium alginate, polyvinyl-pyrrolidine, and/or polyvinyl alcohol, and tableted or encapsulated for conventional administration.
Alternatively, the compounds of this invention may be
20 dissolved in saline, water, polyethylene glycol, propylene glycol, ethanol, corn oil, peanut oil, cottonseed oil, sesame oil, tragacanth gum, and/or various buffers. Other adjuvants and modes of administration are well known in the pharmaceutical art.
25 The carrier or diluent may include time delay material, such as glyceryl monostearate or glyceryl distearate alone or with a wax, or other materials well known in the art.

The pharmaceutical compositions may be made up in a
30 solid form (including granules, powders or suppositories) or in a liquid form (e.g., solutions, suspensions, or emulsions). The pharmaceutical compositions may be subjected to conventional pharmaceutical operations such as sterilization and/or may contain conventional
35 adjuvants, such as preservatives, stabilizers, wetting agents, emulsifiers, buffers etc.

Solid dosage forms for oral administration may include capsules, tablets, pills, powders, and granules. In such solid dosage forms, the active compound may be admixed with at least one inert diluent such as sucrose, 5 lactose, or starch. Such dosage forms may also comprise, as in normal practice, additional substances other than inert diluents, e.g., lubricating agents such as magnesium stearate. In the case of capsules, tablets, and pills, the dosage forms may also comprise 10 buffering agents. Tablets and pills can additionally be prepared with enteric coatings.

Liquid dosage forms for oral administration may include pharmaceutically acceptable emulsions, solutions, suspensions, syrups, and elixirs containing 15 inert diluents commonly used in the art, such as water. Such compositions may also comprise adjuvants, such as wetting, sweetening, flavoring, and perfuming agents.

Compounds of the present invention can possess one or more asymmetric carbon atoms and are thus capable of 20 existing in the form of optical isomers as well as in the form of racemic or non-racemic mixtures thereof. The optical isomers can be obtained by resolution of the racemic mixtures according to conventional processes, e.g., by formation of diastereoisomeric salts, by 25 treatment with an optically active acid or base. Examples of appropriate acids are tartaric, diacetyltaurine, dibenzoyltartaric, ditoluoyltartaric, and camphorsulfonic acid and then separation of the mixture of diastereoisomers by crystallization followed 30 by liberation of the optically active bases from these salts. A different process for separation of optical isomers involves the use of a chiral chromatography column optimally chosen to maximize the separation of the enantiomers. Still another available method involves 35 synthesis of covalent diastereoisomeric molecules by reacting compounds of the invention with an optically pure acid in an activated form or an optically pure

isocyanate. The synthesized diastereoisomers can be separated by conventional means such as chromatography, distillation, crystallization or sublimation, and then hydrolyzed to deliver the enantiomerically pure compound.

5 The optically active compounds of the invention can likewise be obtained by using active starting materials. These isomers may be in the form of a free acid, a free base, an ester or a salt.

The compounds of the present invention can be used
10 in the form of salts derived from inorganic or organic acids. The salts include, but are not limited to, the following: acetate, adipate, alginate, citrate, aspartate, benzoate, benzenesulfonate, bisulfate, butyrate, camphorate, camphorsulfonate, digluconate,
15 cyclopentanepropionate, dodecylsulfate, ethanesulfonate, glucoheptanoate, glycerophosphate, hemisulfate, heptanoate, hexanoate, fumarate, hydrochloride, hydrobromide, hydroiodide, 2-hydroxy-ethanesulfonate, lactate, maleate, methansulfonate, nicotinate, 2-
20 naphthalenesulfonate, oxalate, palmoate, pectinate, persulfate, 2-phenylpropionate, picrate, pivalate, propionate, succinate, tartrate, thiocyanate, tosylate, mesylate, and undecanoate. Also, the basic nitrogen-containing groups can be quaternized with such agents as
25 lower alkyl halides, such as methyl, ethyl, propyl, and butyl chloride, bromides and iodides; dialkyl sulfates like dimethyl, diethyl, dibutyl, and diamyl sulfates, long chain halides such as decyl, lauryl, myristyl and stearyl chlorides, bromides and iodides, aralkyl halides
30 like benzyl and phenethyl bromides, and others. Water or oil-soluble or dispersible products are thereby obtained.

Examples of acids that may be employed to from pharmaceutically acceptable acid addition salts include such inorganic acids as hydrochloric acid, sulphuric acid and phosphoric acid and such organic acids as oxalic acid, maleic acid, succinic acid and citric acid. Other examples include salts with alkali metals or

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alkaline earth metals, such as sodium, potassium, calcium or magnesium or with organic bases.

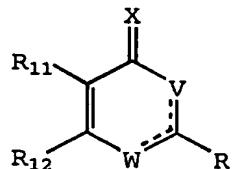
While the compounds of the invention can be administered as the sole active pharmaceutical agent, 5 they can also be used in combination with one or more compounds of the invention or other agents. When administered as a combination, the therapeutic agents can be formulated as separate compositions that are given at the same time or different times, or the 10 therapeutic agents can be given as a single composition.

The foregoing is merely illustrative of the invention and is not intended to limit the invention to the disclosed compounds. Variations and changes which are obvious to one skilled in the art are intended to be 15 within the scope and nature of the invention which are defined in the appended claims.

From the foregoing description, one skilled in the art can easily ascertain the essential characteristics of this invention, and without departing from the spirit 20 and scope thereof, can make various changes and modifications of the invention to adapt it to various usages and conditions.

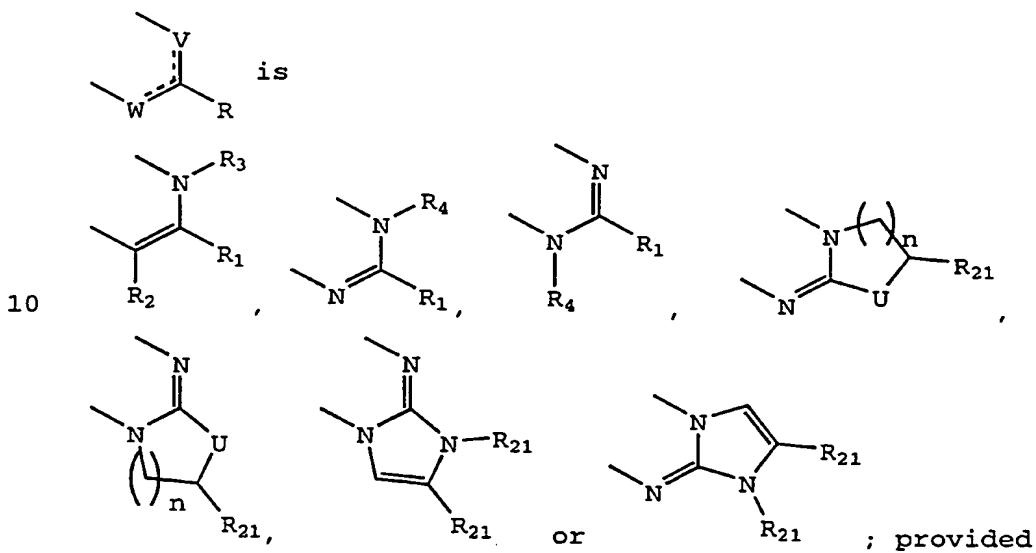
WHAT IS CLAIMED IS:

1. A compound of formula



5 or a pharmaceutically acceptable salt thereof, wherein

X is O, S or NR;:



that the combined total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals in $-VC(R)W-$ is 0-3;

15 U is NR₂₁ or CHR₂₁; and n is an integer of 1-3;

R₁ and R₂ are each independently -Y or -Z-Y, and R₃ and R₄ are each independently -Z-Y; provided that R₄ is other than a hydrogen, substituted-aryl, (substituted-aryl)methyl or (substituted-aryl)ethyl radical, and the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals in each -Y and -Z-Y is 0-3;

wherein each Z is independently a
(1) alkyl, alkenyl or alkynyl radical optionally substituted by (a) 1-3 radicals of amino, alkylamino,
5 dialkylamino, alkanoylamino, alkoxycarbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano or halo, and (b) 1-2 radicals of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino,
10 alkoxycarbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano, halo, alkyl or haloalkyl;
(2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxycarbonylamino, alkylsulfonylamino,
15 hydroxy, alkoxy, alkylthio, cyano, alkyl or haloalkyl; or
(3) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxycarbonylamino, alkylsulfonylamino,
20 hydroxy, alkoxy, alkylthio, cyano, halo, alkyl or haloalkyl;

each Y is independently a
(1) hydrogen radical;
25 (2) halo, cyano or nitro radical;
(3) $-C(O)-R_{20}$, $-C(O)-OR_{21}$, $-C(O)-NR_5R_{21}$ or $-C(NR_5)-NR_5R_{21}$ radical;
(4) $-OR_{21}$, $-O-C(O)-R_{21}$, $-O-C(O)-NR_5R_{21}$ or $-O-C(O)-NR_{22}-S(O)_2-R_{20}$ radical;
30 (5) $-SR_{21}$, $-S(O)-R_{20}$, $-S(O)_2-R_{20}$, $-S(O)_2-NR_5R_{21}$, $-S(O)_2-NR_{22}-C(O)-R_{21}$, $-S(O)_2-NR_{22}-C(O)-OR_{20}$ or $-S(O)_2-NR_{22}-C(O)-NR_5R_{21}$ radical; or
(6) $-NR_5R_{21}$, $-NR_{22}-C(O)-R_{21}$, $-NR_{22}-C(O)-OR_{20}$, $-NR_{22}-C(O)-NR_5R_{21}$, $-NR_{22}-C(NR_5)-NR_5R_{21}$, $-NR_{22}-S(O)_2-R_{20}$ or $-NR_{22}-S(O)_2-NR_5R_{21}$ radical;

wherein each R₅ is independently

- (1) hydrogen radicals;
- (2) alkyl, alkenyl or alkynyl radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, hydroxy, alkoxy, alkylthio, cyano or halo; or
- (3) aryl, heteroaryl, aralkyl, heteroaralkyl, heterocyclyl, heterocyclalkyl, cycloalkyl or cycloalkylalkyl radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, hydroxy, alkoxy, alkylthio, cyano, alkyl or haloalkyl; and

wherein each R₂₀ is independently

- (1) alkyl, alkenyl or alkynyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, N-(alkoxy carbonyl)-N-(alkyl)amino, aminocarbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, cyano, halo or aralkoxy, aralkylthio, aralkylsulfonyl, cycloalkyl, heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, alkanoyl, alkoxy carbonyl, hydroxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, cyano, halo, alkyl or haloalkyl;
- (2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, alkoxy carbonyl, hydroxy, alkoxy, alkylthio, cyano, alkyl or haloalkyl; or
- (3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, alkoxy carbonyl, hydroxy, alkoxy, alkylthio, cyano, halo, azido, alkyl or haloalkyl;

each R₂₁ is independently hydrogen radical or R₂₀;

each R₂₂ is independently

(1) hydrogen radical;

5 (2) alkyl radical optionally substituted by a radical of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, cyano, halo, alkyl or haloalkyl; or
10 (3) heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio,
15 alkylsulfinyl, alkylsulfonyl, cyano, halo, alkyl or haloalkyl; and

each R₂₃ is independently hydrogen or alkyl, or aryl, heteroaryl, aralkyl or heteroaralkyl optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, cyano, halo, alkyl or haloalkyl; and

25 R₁₁ and R₁₂ are each independently an aryl or heteroaryl radical optionally substituted by 1-3 radicals of
(1) R₃₀;
(2) halo or cyano radicals;
30 (3) -C(O)-R₃₀, -C(O)-OR₂₉, -C(O)-NR₃₁R₃₂ or -C(NR₃₁)-NR₃₁R₃₂ radicals;
(4) -OR₂₉, -O-C(O)-R₂₉, -O-C(O)-NR₃₁R₃₂ or -O-C(O)-NR₃₃-S(O)₂-R₃₀ radicals;
(5) -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -S(O)₂-NR₃₃-C(O)-R₃₀, -S(O)₂-NR₃₃-C(O)-NR₃₁R₃₂ radicals; or

(6) $-\text{NR}_{31}\text{R}_{32}$, $-\text{NR}_{33}-\text{C}(\text{O})-\text{R}_{29}$, $-\text{NR}_{33}-\text{C}(\text{O})-\text{OR}_{30}$, $-\text{NR}_{33}-\text{C}(\text{O})-\text{NR}_{31}\text{R}_{32}$, $-\text{NR}_{33}-\text{C}(\text{NR}_{31})-\text{NR}_{31}\text{R}_{32}$, $-\text{NR}_{33}-\text{S}(\text{O})_2-\text{R}_{30}$ or $-\text{NR}_{33}-\text{S}(\text{O})_2-\text{NR}_{31}\text{R}_{32}$ radicals;

provided that (1) R_{11} is other than a 4-pyridyl, 4-pyrimidinyl, 4-quinolyl or 6-isoquinolinyl radical optionally substituted by 1-2 substituents; and (2) the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each of R_{11} and R_{12} is 0-1;

10

wherein each R_{30} is independently

(1) alkyl, alkenyl or alkynyl radicals optionally substituted by 1-3 radicals of $-\text{NR}_{31}\text{R}_{31}$, $-\text{CO}_2\text{R}_{23}$, hydroxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, cyano, halo or aralkoxy, aralkylthio, aralkylsulfonyl, heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, cyano, halo, alkyl or haloalkyl; (2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano, alkyl or haloalkyl; or (3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano, halo, alkyl or haloalkyl;

each R_{29} is independently hydrogen radical or R_{30} ;

35 each R_{31} and R_{32} are each independently
(1) hydrogen radicals;

(2) alkyl radical optionally substituted by an cycloalkyl, aryl, heterocyclyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino,
5 alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano, alkyl or haloalkyl; or
(3) aryl, heteroaryl, heterocyclyl or cycloalkyl radical optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino,
10 alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano, alkyl or haloalkyl; and

wherein each R₃₃ is independently
(1) hydrogen radical; or
15 (2) alkyl radical optionally substituted by a radical of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, alkylamino, dialkylamino, alkanoylamino, alkoxy carbonylamino, alkylsulfonylamino, hydroxy, alkoxy, alkylthio, cyano, alkyl or haloalkyl.
20

2. The compound of Claim 1 or a pharmaceutically acceptable salt thereof, wherein

25 wherein each Z is independently a
(1) C₁-C₈ alkyl, C₂-C₈ alkenyl or C₂-C₈ alkynyl radical optionally substituted by (a) 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄
30 alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano or halo and (b) 1-2 radicals of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
35 alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy,

C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or
(2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or
(3) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

each Y is independently a
(1) hydrogen radical;
(2) halo, cyano or nitro radical;
(3) -C(O)-R₂₀, -C(O)-OR₂₁, -C(O)-NR₅R₂₁ or -C(NR₅)-NR₅R₂₁ radical;
(4) -OR₂₁, -O-C(O)-R₂₁, -O-C(O)-NR₅R₂₁ or -O-C(O)-NR₂₂-S(O)₂-R₂₀ radical;
(5) -SR₂₁, -S(O)-R₂₀, -S(O)₂-R₂₀, -S(O)₂-NR₅R₂₁, -S(O)₂-NR₂₂-C(O)-R₂₁, -S(O)₂-NR₂₂-C(O)-OR₂₀ or -S(O)₂-NR₂₂-C(O)-NR₅R₂₁ radical; or
(6) -NR₅R₂₁, -NR₂₂-C(O)-R₂₁, -NR₂₂-C(O)-OR₂₀, -NR₂₂-C(O)-NR₅R₂₁, -NR₂₂-C(NR₅)-NR₅R₂₁, -NR₂₂-S(O)₂-R₂₀ or -NR₂₂-S(O)₂-NR₅R₂₁ radical;

each R₅ is independently
(1) hydrogen radicals;
(2) C₁-C₈ alkyl, C₂-C₈ alkenyl or C₂-C₈ alkynyl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano or halo; or

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(3) aryl, heteroaryl, aryl-C₁-C₄-alkyl, heteroaryl-C₁-C₄-alkyl, heterocyclyl, heterocyclyl-C₁-C₄-alkyl, C₃-C₈ cycloalkyl or C₃-C₈-cycloalkyl-C₁-C₄-alkyl radicals, optionally substituted by 1-3 radicals of amino, C₁-C₄

5 alkylamino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

each R₂₀ is independently

10 (1) C₁-C₈ alkyl, C₂-C₈ alkenyl or C₂-C₈ alkynyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-C₄ alkyl)amino,

15 aminocarbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, cyano, halo or aryl-C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-alkylsulfonyl, C₃-C₈ cycloalkyl, heterocyclyl, aryl or heteroaryl radicals

20 optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, C₁-C₅ alkanoyl, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄

25 alkylsulfonyl, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

(2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄

30 alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or

(3) aryl or heteroaryl radicals optionally substituted

35 by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄

alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

each R₂₁ is independently hydrogen radical or R₂₀;

each R₂₂ is independently

- 10 (1) hydrogen radical;
- (2) C₁-C₄ alkyl radical optionally substituted by a radical of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or
- (3) heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or aryl, heteroaryl, aryl-C₁-C₄-alkyl or heteroaryl-C₁-C₄-alkyl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

R₁₁ and R₁₂ are each independently an aryl or heteroaryl radical optionally substituted by 1-3 radicals of

(1) R₃₀;

5 (2) halo or cyano radicals;

(3) -C(O)-R₃₀, -C(O)-OR₂₉, -C(O)-NR₃₁R₃₂ or -C(NR₃₁)-NR₃₁R₃₂ radicals;

(4) -OR₂₉, -O-C(O)-R₂₉, -O-C(O)-NR₃₁R₃₂ or -O-C(O)-NR₃₃-S(O)₂-R₃₀ radicals;

10 (5) -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -S(O)₂-NR₃₃-C(O)-R₃₀, -S(O)₂-NR₃₃-C(O)-OR₃₀ or -S(O)₂-NR₃₃-C(O)-NR₃₁R₃₂ radicals; or

(6) -NR₃₁R₃₂, -NR₃₃-C(O)-R₂₉, -NR₃₃-C(O)-OR₃₀, -NR₃₃-C(O)-NR₃₁R₃₂, -NR₃₃-C(NR₃₁)-NR₃₁R₃₂, -NR₃₃-S(O)₂-R₃₀ or -NR₃₃-

15 S(O)₂-NR₃₁R₃₂ radicals;

provided that (1) R₁₁ is other than a 4-pyridyl, 4-pyrimidinyl, 4-quinolyl or 6-isoquinolinyl radical optionally substituted by 1-2 substituents; and (2) the total number of aryl, heteroaryl, cycloalkyl and

20 heterocyclyl radicals substituted on each of R₁₁ and R₁₂ is 0-1;

each R₃₀ is independently

25 (1) C₁-C₄ alkyl, C₂-C₄ alkenyl or C₂-C₄ alkynyl radicals optionally substituted by 1-3 radicals of -NR₃₁R₃₁, -CO₂R₂₃, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, cyano, halo or aryl-

C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-

30 alkylsulfonyl, heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino,

(C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino,

hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄

alkylsulfinyl, C₁-C₄ alkylsulfonyl, cyano, halo, C₁-C₄

35 alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

(2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy,
5 C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or
(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy,
10 C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

each R₂₉ is independently hydrogen radical or R₃₀;

15 each R₃₁ and R₃₂ are each independently
(1) hydrogen radicals;
(2) C₁-C₄ alkyl radical optionally substituted by an C₃-C₈ cycloalkyl, aryl, heterocyclyl or heteroaryl radical
20 optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or
25 (3) aryl, heteroaryl, heterocyclyl or C₃-C₈ cycloalkyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; and
30 alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; and

each R₃₃ is independently

(1) hydrogen radical; or

(2) C₁-C₄ alkyl radical optionally substituted by a radical of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; and

wherein heterocyclyl is a radical of a monocyclic or
10 bicyclic saturated heterocyclic ring system having 5-8 ring members per ring, wherein 1-3 ring members are oxygen, sulfur or nitrogen heteroatoms, which is optionally partially unsaturated or benzo-fused and optionally substituted by 1-2 oxo or thioxo radicals;
15 aryl is a phenyl or naphthyl radical; and heteroaryl is radical of a monocyclic or bicyclic aromatic heterocyclic ring system having 5-6 ring members per ring, wherein 1-3 ring members are oxygen, sulfur or nitrogen heteroatoms, which is optionally benzo-fused or
20 saturated C₃-C₄-carbocyclic-fused.

3. The compound of Claim 2 or a pharmaceutically acceptable salt thereof, wherein
25 each Z is independently a
(1) C₁-C₈ alkyl, C₂-C₈ alkenyl or C₂-C₈ alkynyl radical optionally substituted by (a) 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo and (b) 1-2 radicals of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy,

C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

(2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or

(3) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

15 each R₅ is independently

(1) hydrogen radicals;

(2) C₁-C₄ alkyl, C₂-C₅ alkenyl or C₂-C₅ alkynyl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo; or

(3) aryl, heteroaryl, aryl-C₁-C₄-alkyl, heteroaryl-C₁-C₄-alkyl, heterocyclyl, heterocyclyl-C₁-C₄-alkyl, C₃-C₈ cycloalkyl or C₃-C₈-cycloalkyl-C₁-C₄-alkyl radicals

20 optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

25 each R₂₀ is independently

(1) C₁-C₈ alkyl, C₂-C₅ alkenyl or C₂-C₅ alkynyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-C₄ alkyl)amino,

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aminocarbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo or aryl-C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-alkylsulfonyl, C₃-C₈ cycloalkyl,

5 heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, C₁-C₅ alkanoyl, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy,

10 C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

(2) heterocyclyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or

15 (3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

20 by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

25 cyano, halo, azido, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

each R₂₁ is independently hydrogen radical or R₂₀;

each R₃₀ is independently

30 (1) C₁-C₄ alkyl radical optionally substituted by 1-3 radicals of

(a) -NR₃₁R₃₁;

(b) C₁-C₄ alkoxy-carbonyl or phenoxy carbonyl or phenylmethoxy carbonyl optionally substituted by 1-3

35 radicals of amino, alkylamino, di-(C₁-C₄-alkyl)amino,

C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl; or

5 (c) hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, or phenyl-C₁-C₄-alkoxy, phenyl-C₁-C₄-alkylthio, heterocyclyl, phenyl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals;

10 (2) C₁-C₄ haloalkyl of 1-3 halo radical; or

(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

15 20 each R₂₉ is independently hydrogen radical or R₃₀;

each R₃₁ is independently

(1) hydrogen radicals; or

25 (2) C₁-C₄ alkyl radical optionally substituted by an phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or trifluoromethyl radicals; and

30 each R₃₂ is independently

(1) hydrogen radicals;

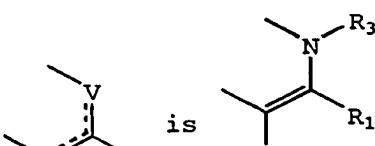
(2) C₁-C₄ alkyl radical optionally substituted by an C₃-C₆ cycloalkyl, aryl, heterocyclyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; or

(3) aryl, heteroaryl, heterocyclyl or C₃-C₆ cycloalkyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₄ haloalkyl of 1-3 halo radicals; and

each R₃₃ is independently hydrogen or C₁-C₄ alkyl radical.

4. The compound of Claim 3 or a pharmaceutically acceptable salt thereof, wherein

X is O or S;

25  ; provided that the combined total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals in -VC(R)W- is 0-2;

wherein R₁ is -Y or -Z-Y, provided that (1) the total
30 number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals in R₁ is 0-3;

Z is a

(1) C₁-C₈ alkyl or C₂-C₈ alkenyl radical optionally substituted by (a) 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo and (b) 1-2 radicals of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

(2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, di-(C₁-C₄ alkyl)amino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl radicals; or

(3) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

Y is a

(1) hydrogen radical;

(2) halo radical;

(3) -C(O)-R₂₀, -C(O)-OR₂₁, -C(O)-NR₅R₂₁ or -C(NR₅)-NR₅R₂₁ radical;

(4) -OR₂₁, -O-C(O)-R₂₁ or -O-C(O)-NR₅R₂₁ radical;

(5) -SR₂₁, -S(O)-R₂₀, -S(O)₂-R₂₀ or -S(O)₂-NR₅R₂₁ radical;

or

(6) -NR₅R₂₁, -NR₂₂-C(O)-R₂₁, -NR₂₂-C(O)-OR₂₀, -NR₂₂-C(O)-NR₅R₂₁, -NR₂₂-C(NR₅)-NR₅R₂₁, -NR₂₂-S(O)₂-R₂₀ or -NR₂₂-S(O)₂-NR₅R₂₁ radical;

each R₅ is independently

- (1) hydrogen radicals;
- (2) C₁-C₄ alkyl or C₂-C₅ alkenyl radicals optionally substituted by 1-3 radicals of amino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo; or
- (3) phenyl-C₁-C₂-alkyl, heteroaryl-C₁-C₂-alkyl, heterocyclyl-C₁-C₂-alkyl or C₃-C₆-cycloalkyl-C₁-C₂-alkyl radicals optionally substituted by 1-3 radicals of amino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

each R₂₀ is independently

- (1) C₁-C₈ alkyl or C₂-C₅ alkenyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-C₄ alkyl)amino, aminocarbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo or aryl-C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-alkylsulfonyl, C₃-C₆ cycloalkyl, heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, C₁-C₅ alkanoyl, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;
- (2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl; or

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(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

each R₂₁ is independently hydrogen radical or R₂₀;

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each R₂₂ is independently
(1) hydrogen radical; or
(2) C₁-C₄ alkyl radical optionally substituted by a radical of phenyl or heteroaryl optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or phenyl, heteroaryl, phenyl-C₁-C₂-alkyl or heteroaryl-C₁-C₂-alkyl optionally substituted by 1-3 radicals of amino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

R₂ is a radical of hydrogen, C₁-C₄ alkyl, halo, cyano, hydroxy, C₁-C₄ alkoxy, C₁-C₂ haloalkoxy of 1-3 halo radicals, C₁-C₄ alkylthio, amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino or C₁-C₂ haloalkyl of 1-3 halo radicals;

R₃ is a hydrogen radical or

(1) C₁-C₈ alkyl or C₂-C₈ alkenyl radical optionally substituted by (a) 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, 5 hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo, and (b) 1-2 radicals of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, 10 hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals; or (2) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ 15 alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;

R₁₁ and R₁₂ are each independently an aryl or heteroaryl 20 radical optionally substituted by 1-2 radicals of

(1) R₃₀;
(2) halo or cyano radicals;
(3) -C(O)-R₃₀, -C(O)-OR₂₉, -C(O)-NR₃₁R₃₂ or -C(NR₃₁)-NR₃₁R₃₂ radicals; or
25 (4) -OR₂₉, -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -NR₃₁R₃₂, -NR₃₃-C(O)-R₂₉ or -NR₃₃-C(O)-OR₃₀ radicals; provided that (1) R₁₁ is other than a 4-pyridyl, 4-pyrimidinyl, 4-quinolyl or 6-isoquinolinyl radical 30 optionally substituted by 1-2 substituents; and (2) the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each of R₁₁ and R₁₂ is 0-1;

each R₃₀ is independently
35 (1) C₁-C₄ alkyl radical optionally substituted by

- (a) amino, C₁-C₄ alkylamino or di-(C₁-C₄-alkyl)amino radicals; or
- (b) hydroxy, C₁-C₄ alkoxy, heterocyclyl, phenyl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

5. 10 (2) C₁-C₂ haloalkyl of 1-3 halo radical; or
(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

15. 20 each R₂₉ is independently hydrogen radical or R₃₀;

each R₃₁ is independently hydrogen or C₁-C₄ alkyl radicals; and

each R₃₂ is independently

- (1) hydrogen radicals;
- (2) C₁-C₄ alkyl radical optionally substituted by phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl or trifluoromethyl radicals; or
- (3) phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl or trifluoromethyl radicals; and

each R₃₃ is independently hydrogen or methyl radical;
and

5 wherein heterocycll is a radical of a monocyclic
saturated heterocyclic ring system having 5-6 ring
members, wherein 1-3 ring members are oxygen, sulfur or
nitrogen heteroatoms, which is optionally benzo-fused
and optionally substituted by 1-2 oxo or thioxo
radicals; aryl is a phenyl or naphthyl radical; and
10 heteroaryl is radical of a monocyclic aromatic
heterocyclic ring system having 5-6 ring members,
wherein 1-3 ring members are oxygen, sulfur or nitrogen
heteroatoms, which is optionally benzo-fused or
15 saturated C₃-C₄-carbocyclic-fused.

5. The compound of Claim 4 or a pharmaceutically
acceptable salt thereof, wherein

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Z is a

(1) C₁-C₄ alkyl or C₂-C₅ alkenyl radical optionally
substituted by (a) 1-3 radicals of amino, di-(C₁-C₂
alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
25 alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂
alkylthio or halo and (b) 1-2 radicals of heterocycll,
aryl or heteroaryl optionally substituted by 1-3
radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₂
alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
30 alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄
alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl
radicals;
(2) heterocycll radical optionally substituted by 1-2
radicals of amino, di-(C₁-C₂ alkyl)amino, (C₁-C₄
35 alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂
alkylthio or C₁-C₄ alkyl radicals; or

(3) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or 5 trifluoromethyl radicals;

each R₅ is independently

(1) hydrogen radical;
(2) C₁-C₄ alkyl radical optionally substituted by 1-3
10 radicals of amino, di-(C₁-C₂-alkyl)amino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio or halo; or
(3) phenyl-C₁-C₂-alkyl, heteroaryl-C₁-C₂-alkyl,
heterocycl-C₁-C₂-alkyl or C₃-C₆-cycloalkyl-C₁-C₂-alkyl
15 radicals optionally substituted by 1-3 radicals of
amino, di-(C₁-C₂-alkyl)amino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, methoxy, methylthio, cyano, C₁-C₄ alkyl or
trifluoromethyl radicals;

each R₂₂ is independently hydrogen or C₁-C₄ alkyl
20 radical;

each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or
phenyl, heteroaryl, phenyl-C₁-C₂-alkyl or heteroaryl-C₁-C₂-alkyl optionally substituted by 1-3 radicals of
25 amino, di-(C₁-C₂ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl
radicals;

30 R₃ is a hydrogen radical or
(1) C₁-C₈ alkyl radical optionally substituted by 1-2
radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy or aryl or heteroaryl
optionally substituted by 1-3 radicals of amino, C₁-C₄
35 alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄

alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals; or
(2) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;

R₁₁ is an aryl radical and R₁₂ is a heteroaryl radical,
10 wherein the aryl and heteroaryl radicals are optionally substituted by 1-2 radicals of

(1) R₃₀;
(2) halo or cyano radicals;
(3) -C(O)-R₃₀, -C(O)-OR₂₉, -C(O)-NR₃₁R₃₂ or -C(NR₃₁)-NR₃₁R₃₂ radicals; or
(4) -OR₂₉, -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -NR₃₁R₃₂ or -NR₃₃-C(O)-R₂₉ radicals;
provided that the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each
20 of R₁₁ and R₁₂ is 0-1;

each R₃₀ is independently
(1) C₁-C₄ alkyl radical optionally substituted by a phenyl or heteroaryl radical optionally substituted by
25 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, hydroxy, C₁-C₂ alkoxy, halo, C₁-C₄ alkyl or trifluoromethyl radicals;
(2) trifluoromethyl radical; or
(3) aryl or heteroaryl radicals optionally substituted
30 by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, hydroxy, C₁-C₂ alkoxy, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

each R₂₉ is independently hydrogen radical or R₃₀; and
35 each R₃₂ is independently

(1) hydrogen radicals;
(2) C₁-C₄ alkyl radical or C₁-C₂ alkyl radical substituted by phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, hydroxy, C₁-C₂ alkoxy, C₁-C₄ alkyl or trifluoromethyl radicals; or
(3) phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, hydroxy, C₁-C₂ alkoxy, C₁-C₄ alkyl or trifluoromethyl radicals; and
wherein heterocyclyl is a radical of a monocyclic saturated heterocyclic ring system having 5-6 ring members, wherein 1-2 ring members are oxygen, sulfur or nitrogen heteroatoms, which is optionally benzo-fused and optionally substituted by 1-2 oxo or thioxo radicals; aryl is a phenyl or naphthyl radical; and heteroaryl is radical of a monocyclic aromatic heterocyclic ring system having 5-6 ring members, wherein 1-2 ring members are oxygen, sulfur or nitrogen heteroatoms, which is optionally benzo-fused.

6. The compound of Claim 5 or a pharmaceutically acceptable salt thereof, wherein

wherein R₁ is -Y or -Z-Y, provided that (1) the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals in R₁ is 0-2;

Z is a
(1) C₁-C₄ alkyl or C₂-C₅ alkenyl radical optionally substituted by (a) 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio or halo and (b) 1-2 radicals of aryl or heteroaryl optionally substituted by 1-2

radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals; or

5 (2) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

10

Y is a

- (1) hydrogen radical;
- (2) -C(O)-R₂₀, -C(O)-OR₂₁ or -C(O)-NR₅R₂₁ radical;
- (3) -OR₂₁, -SR₂₁, -S(O)-R₂₀, -S(O)₂-R₂₀ or -S(O)₂-NR₅R₂₁
- 15 (4) -NR₅R₂₁, -NR₂₂-C(O)-R₂₁, -NR₂₂-C(O)-OR₂₀, -NR₂₂-C(O)-NR₅R₂₁, -NR₂₂-S(O)₂-R₂₀ or -NR₂₂-S(O)₂-NR₅R₂₁ radical;

each R₅ is independently

20 (1) hydrogen radical;

- (2) C₁-C₄ alkyl radical optionally substituted by 1-3 halo radicals; or
- (3) phenyl-C₁-C₂-alkyl or heteroaryl-C₁-C₂-alkyl, radicals optionally substituted by 1-3 radicals of

25 amino, dimethylamino, hydroxy, methoxy, methylthio, methyl or trifluoromethyl radicals;

each R₂₀ is independently

- (1) C₁-C₈ alkyl or C₂-C₅ alkenyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-C₄ alkyl)amino, aminocarbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo or aryl-C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-alkylsulfonyl, C₃-C₆ cycloalkyl,

heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, C₁-C₅

5 alkanoyl, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

(2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, di-(C₁-C₄ alkyl)amino, (C₁-C₄ alkoxy)carbonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl; or

(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, acetamido, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or trifluoromethyl radicals;

each R₂₁ is independently hydrogen radical or R₂₀;

20 each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or phenyl-C₁-C₂-alkyl or heteroaryl-C₁-C₂-alkyl optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, (C₁-C₄ alkoxy)carbonylamino,

25 hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

R₂ is a radical of hydrogen, C₁-C₄ alkyl, halo, cyano, hydroxy, C₁-C₄ alkoxy, trifluoromethoxy or

30 trifluoromethyl;

R₃ is a hydrogen radical or C₁-C₈ alkyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy

35 or aryl or heteroaryl optionally substituted by 1-3

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radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;

5 R₁₁ is an aryl radical and R₁₂ is a heteroaryl radical, wherein the aryl and heteroaryl radicals are optionally substituted by 1-2 radicals of

- (1) R₃₀;
- (2) halo or cyano radicals; or
- 10 (3) -C(O)-NR₃₁R₃₂, -OR₂₉, -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -NR₃₁R₃₂ or -NR₃₃-C(O)-R₂₉ radicals; provided that the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each of R₁₁ and R₁₂ is 0-1;

15 each R₃₀ is independently

- (1) C₁-C₄ alkyl radical optionally substituted by a phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;
- (2) trifluoromethyl radical; or
- (3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;

each R₂₉ is independently hydrogen radical or R₃₀;

30 each R₃₁ is independently hydrogen, methyl or ethyl radicals; and

each R₃₂ is independently

- (1) hydrogen radicals;
- 35 (2) C₁-C₄ alkyl radical or C₁-C₂ alkyl radical substituted by phenyl or heteroaryl radical optionally

substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals; or

5 (3) phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals.

7. The compound of Claim 6 or a pharmaceutically
10 acceptable salt thereof, wherein

R₃ is a radical of hydrogen or C₁-C₄ alkyl;

15 R₁₁ is an aryl radical optionally substituted by 1-2
radicals of

- (1) R₃₀;
- (2) halo or cyano radicals; or
- (3) -C(O)-NR₃₁R₃₂, -OR₂₉, -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -NR₃₁R₃₂ or -NR₃₃-C(O)-R₂₉ radicals; and

20 R₁₂ is a heteroaryl radical optionally substituted by 1-2
radicals of

- (1) R₃₀;
- (2) halo or cyano radicals; or
- (3) -C(O)-NR₃₁R₃₂, -OR₂₉, -SR₂₉, -NR₃₁R₃₂ or -NR₃₃-C(O)-R₂₉ radicals;

provided that the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each of R₁₁ and R₁₂ is 0-1;

30 R₃₀ is independently

- (1) C₁-C₄ alkyl radical optionally substituted by a phenyl or heteroaryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;

(2) trifluoromethyl radical; or
(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl
5 radicals;

R₂₉ is an aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or
10 trifluoromethyl radicals; and

R₃₂ is independently
(1) hydrogen or C₁-C₄ alkyl radical; or
(2) phenyl or heteroaryl radical optionally substituted
15 by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals.

8. The compound of Claim 7 or a pharmaceutically
20 acceptable salt thereof, wherein

wherein R₁ is -Y or -Z-Y, provided that (1) the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals in R₁ is 0-1;
25 Z is a C₁-C₄ alkyl radical optionally substituted by 1-2 radicals of amino, di-(C₁-C₂ alkyl)amino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, halo, or aryl or heteroaryl optionally substituted by 1-2 radicals of hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;
30 each R₅ is independently hydrogen or C₁-C₄ alkyl
35 radical;

each R₂₀ is independently

(1) C₁-C₈ alkyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄

5 alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-C₄ alkyl)amino, aminocarbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo or C₃-C₆ cycloalkyl, heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-10 2 radicals of amino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

15 (2) heterocyclyl radical optionally substituted by 1-2 radicals of (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl; or
(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of (C₁-C₄ alkoxy)carbonyl, amino, C₁-C₄ 20 alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or trifluoromethyl radicals;

each R₂₁ is independently hydrogen radical or R₂₀;

25 each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or phenyl-C₁-C₂-alkyl optionally substituted by 1-2 radicals of hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

30 R₂ is a hydrogen radical;

R₃ is a hydrogen, methyl or ethyl radical;

R₁₁ is an aryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methylthio, methylsulfinyl, methylsulfonyl, aminocarbonyl, methyl or trifluoromethyl
5 radicals; and

R₁₂ is a heteroaryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methyl or trifluoromethyl
10 radicals.

9. The compound of Claim 8 or a pharmaceutically acceptable salt thereof, wherein
15

Z is C₁-C₄ alkyl radical optionally substituted by 1-2 radicals of amino, t-butoxycarbonylamino, dimethylamino, hydroxy, methoxy, methylthio or halo radicals;

20 Y is a

- (1) hydrogen radical;
- (2) -C(O)-R₂₀, -C(O)-OR₂₁ or -C(O)-NR₅R₂₁ radical;
- (3) -OR₂₁, -SR₂₁, -S(O)-R₂₀, -S(O)₂-R₂₀ or -S(O)₂-NR₅R₂₁ radical; or

25 (4) -NR₅R₂₁, -NR₂₂-C(O)-R₂₁ or -NR₂₂-S(O)₂-R₂₀ radical;

R₅ is a hydrogen radical;

each R₂₀ is independently

30 (1) C₁-C₆ alkyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, methylamino, dimethylamino, t-butoxycarbonylamino, N-((t-butoxy)carbonyl)-N-(methyl)amino, aminocarbonylamino, hydroxy, butoxy, methoxy, butylthio, methylthio, methylsulfinyl,
35 methylsulfonyl, halo or C₅-C₆ cycloalkyl, heterocyclyl, phenyl or heteroaryl radicals optionally substituted by

1-2 radicals of amino, dimethylamino, acetamino, hydroxy, methoxy, methylthio, halo, methyl or trifluoromethyl radicals;

(2) heterocyclyl radical optionally substituted by 1-2 radicals of t-butoxycarbonyl, hydroxy, or C₁-C₄ alkyl; or

(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, hydroxy, methoxy, methylthio, halo, methyl or trifluoromethyl radicals;

each R₂₁ is independently hydrogen radical or R₂₀;

each R₂₂ is independently hydrogen or methyl radical;

each R₂₃ is independently hydrogen or C₁-C₄ alkyl radicals;

R₁₁ is an unsubstituted phenyl or naphthyl radical or a phenyl radical substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methylthio, methylsulfinyl, methylsulfonyl, aminocarbonyl, methyl or trifluoromethyl radicals; and

R₁₂ is a 4-pyridyl, 4-quinolinyl, 4-imidazolyl or 4-pyrimidinyl radical optionally substituted by a radical of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methyl or trifluoromethyl radicals.

10. The compound of Claim 9 or a pharmaceutically acceptable salt thereof, wherein

Y is a

(1) -C(O)-R₂₀ or -C(O)-NR₅R₂₁ radical;

(2) -OR₂₁, -SR₂₁, -S(O)-R₂₀, -S(O)₂-R₂₀ or -S(O)₂-NR₅R₂₁ radical; or

(3) $-\text{NR}_5\text{R}_{21}$, $-\text{NR}_{22}-\text{C}(\text{O})-\text{R}_{21}$ or $-\text{NR}_{22}-\text{S}(\text{O})_2-\text{R}_{20}$ radical;

each R_{20} is independently

(1) $\text{C}_1\text{-C}_6$ alkyl radicals optionally substituted by 1-3
5 radicals of $-\text{CO}_2\text{R}_{23}$, amino, methylamino, dimethylamino,
 t -butoxycarbonylamino, $\text{N}-((t\text{-butoxy)carbonyl})-\text{N}$ -
(methyl)amino, aminocarbonylamino, hydroxy, butoxy,
methoxy, butylthio, methylthio, methylsulfinyl,
methylsulfonyl, halo or $\text{C}_5\text{-C}_6$ cycloalkyl, heterocyclyl,
10 phenyl or heteroaryl radicals optionally substituted by
1-2 radicals of amino, dimethylamino, acetamino,
hydroxy, methoxy, methylthio, halo, methyl or
trifluoromethyl radicals;
(2) heterocyclyl radical optionally substituted by t -
15 butoxycarbonyl; or
(3) aryl or heteroaryl radicals optionally substituted
by 1-2 radicals of amino, dimethylamino, hydroxy,
methoxy, methylthio, halo, methyl or trifluoromethyl
radicals; and

20 each R_{21} is independently hydrogen radical or R_{20} .

11. The compound of Claim 10 or a pharmaceutically
25 acceptable salt thereof, wherein

Y is a $-\text{OR}_{21}$, $-\text{SR}_{21}$ or $-\text{NR}_5\text{R}_{21}$ radical;

each R_{20} is independently

30 (1) $\text{C}_1\text{-C}_6$ alkyl radicals optionally substituted by 1-3
radicals of amino, methylamino, dimethylamino, hydroxy
or phenyl or heteroaryl radicals optionally substituted
by 1-2 radicals of amino, dimethylamino, hydroxy,
methoxy, methylthio, halo, methyl or trifluoromethyl
35 radicals;
(2) heterocyclyl radical; or

(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, hydroxy, methoxy, methylthio, halo, methyl or trifluoromethyl radicals;

5

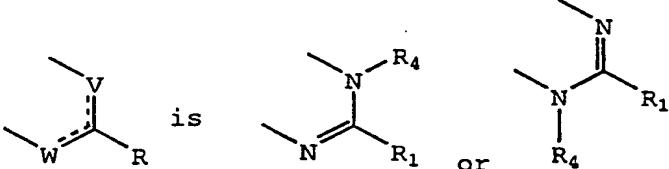
each R₂₁ is independently hydrogen radical or R₂₀;

R₁₁ is an unsubstituted phenyl radical or a phenyl radical substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methylthio, methylsulfonyl, methyl or trifluoromethyl radicals; and

R₁₂ is a 4-pyridyl radical optionally substituted by a radical of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methyl or trifluoromethyl radicals.

20 12. The compound of Claim 3 or a pharmaceutically acceptable salt thereof, wherein

X is O or S;

25  is ; provided that the combined total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals in -VC(R)W- is 0-2;

wherein R₁ is -Y or -Z-Y, provided that (1) the total 30 number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals in R₁ is 0-3;

Z is a

(1) C₁-C₈ alkyl or C₂-C₈ alkenyl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄)alkoxy carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, or heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄)alkoxy carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

(2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, di-(C₁-C₄ alkyl)amino, (C₁-C₄)alkoxy carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl radicals; or

(3) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄)alkylamino, C₁-C₅ alkanoylamino, (C₁-C₄)alkoxy carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

Y is a

(1) hydrogen radical;

(2) halo radical;

(3) -C(O)-R₂₀, -C(O)-OR₂₁, -C(O)-NR₅R₂₁ or -C(NR₅)-NR₅R₂₁ radical;

(4) -OR₂₁, -O-C(O)-R₂₁ or -O-C(O)-NR₅R₂₁ radical;

(5) -SR₂₁, -S(O)-R₂₀, -S(O)₂-R₂₀ or -S(O)₂-NR₅R₂₁ radical;

or

(6) -NR₅R₂₁, -NR₂₂-C(O)-R₂₁, -NR₂₂-C(O)-OR₂₀, -NR₂₂-C(O)-NR₅R₂₁, -NR₂₂-C(NR₅)-NR₅R₂₁, -NR₂₂-S(O)₂-R₂₀ or -NR₂₂-S(O)₂-NR₅R₂₁ radical;

each R₅ is independently

(1) hydrogen radicals;

(2) C₁-C₄ alkyl or C₂-C₅ alkenyl radicals optionally substituted by 1-3 radicals of amino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo; or

5 (3) phenyl-C₁-C₂-alkyl, heteroaryl-C₁-C₂-alkyl, heterocyclyl-C₁-C₂-alkyl or C₃-C₆-cycloalkyl-C₁-C₂-alkyl radicals optionally substituted by 1-3 radicals of amino, di-(C₁-C₄-alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-10 3 halo radicals;

each R₂₀ is independently

(1) C₁-C₈ alkyl or C₂-C₅ alkenyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-C₄ alkyl)amino, aminocarbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo or aryl-C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-alkylsulfonyl, C₃-C₆ cycloalkyl, heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, C₁-C₅ alkanoyl, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

20 (2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl; or

25 (3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄

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alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

5

each R₂₁ is independently hydrogen radical or R₂₀;

each R₂₂ is independently

(1) hydrogen radical; or

10 (2) C₁-C₄ alkyl radical optionally substituted by a radical of phenyl or heteroaryl optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or
15 C₁-C₂ haloalkyl of 1-3 halo radicals;

each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or phenyl, heteroaryl, phenyl-C₁-C₂-alkyl or heteroaryl-C₁-C₂-alkyl optionally substituted by 1-3 radicals of
20 amino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;

25 R₄ is

(1) C₁-C₈ alkyl or C₂-C₈ alkenyl radical optionally substituted by (a) 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino,
30 hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or halo, and (b) 1-2 radicals of heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino,

hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals; or
(2) heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;

5 R₁₁ and R₁₂ are each independently an aryl or heteroaryl radical optionally substituted by 1-2 radicals of
(1) R₃₀;
(2) halo or cyano radicals;
(3) -C(O)-R₃₀, -C(O)-OR₂₉, -C(O)-NR₃₁R₃₂ or -C(NR₃₁)-

10 NR₃₁R₃₂ radicals; or
(4) -OR₂₉, -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂,
-NR₃₁R₃₂, -NR₃₃-C(O)-R₂₉ or -NR₃₃-C(O)-OR₃₀ radicals;
provided that (1) R₁₁ is other than a 4-pyridyl, 4-pyrimidinyl, 4-quinolyl or 6-isouquinolinyl radical
15 20 optionally substituted by 1-2 substituents; and (2) the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each of R₁₁ and R₁₂ is 0-1;

25 each R₃₀ is independently
(1) C₁-C₄ alkyl radical optionally substituted by
(a) amino, C₁-C₄ alkylamino or di-(C₁-C₄-alkyl)amino radicals; or
(b) hydroxy, C₁-C₄ alkoxy, heterocyclyl, phenyl or
30 heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl
35 radicals;

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(2) C₁-C₂ haloalkyl of 1-3 halo radical; or
(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

each R₂₉ is independently hydrogen radical or R₃₀;

each R₃₁ is independently hydrogen or C₁-C₄ alkyl radicals; and

each R₃₂ is independently
(1) hydrogen radicals;
(2) C₁-C₄ alkyl radical optionally substituted by phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl or trifluoromethyl radicals; or
(3) phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl or trifluoromethyl radicals; and

each R₃₃ is independently hydrogen or methyl radical; and

wherein heterocyclyl is a radical of a monocyclic saturated heterocyclic ring system having 5-6 ring members, wherein 1-3 ring members are oxygen, sulfur or nitrogen heteroatoms, which is optionally benzo-fused and optionally substituted by 1-2 oxo or thioxo radicals; aryl is a phenyl or naphthyl radical; and

heteroaryl is radical of a monocyclic aromatic heterocyclic ring system having 5-6 ring members, wherein 1-3 ring members are oxygen, sulfur or nitrogen heteroatoms, which is optionally benzo-fused or
5 saturated C₃-C₄-carbocyclic-fused.

13. The compound of Claim 12 or a pharmaceutically acceptable salt thereof, wherein

10

Z is a

- (1) C₁-C₄ alkyl or C₂-C₅ alkenyl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ 15 alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, halo, or heterocyclyl, aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₂ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ 20 alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;
- (2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, di-(C₁-C₂ alkyl)amino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ 25 alkylthio or C₁-C₄ alkyl radicals; or
- (3) aryl or heteroaryl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or 30 trifluoromethyl radicals;

each R₅ is independently

- (1) hydrogen radical;
- (2) C₁-C₄ alkyl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂-alkyl)amino, hydroxy, C₁-C₂ 35 alkoxy, C₁-C₂ alkylthio or halo; or

(3) phenyl-C₁-C₂-alkyl, heteroaryl-C₁-C₂-alkyl, heterocycll-C₁-C₂-alkyl or C₃-C₆-cycloalkyl-C₁-C₂-alkyl radicals optionally substituted by 1-3 radicals of amino, di-(C₁-C₂-alkyl)amino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, methoxy, methylthio, cyano, C₁-C₄ alkyl or trifluoromethyl radicals;

each R₂₂ is independently hydrogen or C₁-C₄ alkyl radical;

each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or phenyl, heteroaryl, phenyl-C₁-C₂-alkyl or heteroaryl-C₁-C₂-alkyl optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

R₄ is

(1) C₁-C₈ alkyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy or aryl or heteroaryl optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals; or
(2) heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo.

C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;

R₁₁ is an aryl radical and R₁₂ is a heteroaryl radical, wherein the aryl and heteroaryl radicals are optionally substituted by 1-2 radicals of

(1) R_{30} ;

(2) halo or cyano radicals;

(3) $-C(O)-R_{30}$, $-C(O)-OR_{29}$, $-C(O)-NR_{31}R_{32}$ or $-C(NR_{31})-$
5 $NR_{31}R_{32}$ radicals; or

(4) $-OR_{29}$, $-SR_{29}$, $-S(O)-R_{30}$, $-S(O)_2-R_{30}$, $-S(O)_2-NR_{31}R_{32}$,
 $-NR_{31}R_{32}$ or $-NR_{33}-C(O)-R_{29}$ radicals;
provided that the total number of aryl, heteroaryl,
cycloalkyl and heterocyclyl radicals substituted on each
of R_{11} and R_{12} is 0-1;

10 each R_{30} is independently

(1) C_1-C_4 alkyl radical optionally substituted by a
phenyl or heteroaryl radical optionally substituted by
1-3 radicals of amino, di- $(C_1-C_2$ alkyl)amino, acetamido,
15 hydroxy, C_1-C_2 alkoxy, halo, C_1-C_4 alkyl or
trifluoromethyl radicals;

(2) trifluoromethyl radical; or

(3) aryl or heteroaryl radicals optionally substituted
by 1-3 radicals of amino, di- $(C_1-C_2$ alkyl)amino,
20 acetamido, hydroxy, C_1-C_2 alkoxy, halo, C_1-C_4 alkyl or
trifluoromethyl radicals;

each R_{29} is independently hydrogen radical or R_{30} ; and

25 each R_{32} is independently

(1) hydrogen radicals;

(2) C_1-C_4 alkyl radical or C_1-C_2 alkyl radical
substituted by phenyl or heteroaryl radical optionally
substituted by 1-3 radicals of amino, di- $(C_1-C_2$
30 alkyl)amino, acetamido, hydroxy, C_1-C_2 alkoxy, C_1-C_4
alkyl or trifluoromethyl radicals; or

(3) phenyl or heteroaryl radical optionally substituted
by 1-3 radicals of amino, di- $(C_1-C_2$ alkyl)amino,
acetamido, hydroxy, C_1-C_2 alkoxy, C_1-C_4 alkyl or
35 trifluoromethyl radicals; and

wherein heterocyclyl is a radical of a monocyclic saturated heterocyclic ring system having 5-6 ring members, wherein 1-2 ring members are oxygen, sulfur or nitrogen heteroatoms, which is optionally benzo-fused
5 and optionally substituted by 1-2 oxo or thioxo radicals; aryl is a phenyl or naphthyl radical; and heteroaryl is radical of a monocyclic aromatic heterocyclic ring system having 5-6 ring members, wherein 1-2 ring members are oxygen, sulfur or nitrogen
10 heteroatoms, which is optionally benzo-fused.

14. The compound of Claim 13 or a pharmaceutically acceptable salt thereof, wherein

15 wherein R₁ is -Y or -Z-Y, provided that (1) the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals in R₁ is 0-2;

20 Z is a

(1) C₁-C₄ alkyl or C₂-C₅ alkenyl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, halo, or aryl or heteroaryl

25 optionally substituted by 1-2 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals; or

(2) aryl or heteroaryl radical optionally substituted by
30 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

35 Y is a

(1) hydrogen radical;

(2) $-C(O)-R_{20}$, $-C(O)-OR_{21}$ or $-C(O)-NR_5R_{21}$ radical;
(3) $-OR_{21}$, $-SR_{21}$, $-S(O)-R_{20}$, $-S(O)_2-R_{20}$ or $-S(O)_2-NR_5R_{21}$ radical; or
(4) $-NR_5R_{21}$, $-NR_{22}-C(O)-R_{21}$, $-NR_{22}-C(O)-OR_{20}$, $-NR_{22}-C(O)-NR_5R_{21}$,
5 $-NR_{22}-S(O)_2-R_{20}$ or $-NR_{22}-S(O)_2-NR_5R_{21}$ radical;

each R_5 is independently
(1) hydrogen radical;
(2) C_1-C_4 alkyl radical optionally substituted by 1-3
10 halo radicals; or
(3) phenyl- C_1-C_2 -alkyl or heteroaryl- C_1-C_2 -alkyl,
radicals optionally substituted by 1-3 radicals of
amino, dimethylamino, hydroxy, methoxy, methylthio,
methyl or trifluoromethyl radicals;

15 each R_{20} is independently
(1) C_1-C_8 alkyl or C_2-C_5 alkenyl radicals optionally
substituted by 1-3 radicals of $-CO_2R_{23}$, amino, C_1-C_4
alkylamino, di-(C_1-C_4 alkyl)amino, C_1-C_5 alkanoylamino,
20 (C_1-C_4 alkoxy)carbonylamino, $N-((C_1-C_4$ alkoxy)carbonyl)-
 $N-(C_1-C_4$ alkyl)amino, aminocarbonylamino, hydroxy, C_1-C_4
alkoxy, C_1-C_4 alkylthio, C_1-C_4 alkylsulfinyl, C_1-C_4
alkylsulfonyl, halo or aryl- C_1-C_4 -alkoxy, aryl- C_1-C_4 -
alkylthio, aryl- C_1-C_4 -alkylsulfonyl, C_3-C_6 cycloalkyl,
25 heterocyclyl, aryl or heteroaryl radicals optionally
substituted by 1-3 radicals of amino, C_1-C_4 alkylamino,
di-(C_1-C_4 alkyl)amino, C_1-C_5 alkanoylamino, (C_1-C_4
alkoxy)carbonylamino, C_1-C_4 alkylsulfonylamino, C_1-C_5
alkanoyl, (C_1-C_4 alkoxy)carbonyl, hydroxy, C_1-C_4 alkoxy,
30 C_1-C_4 alkylthio, cyano, halo, C_1-C_4 alkyl or C_1-C_2
haloalkyl of 1-3 halo radicals;
(2) heterocyclyl radical optionally substituted by 1-2
radicals of amino, di-(C_1-C_4 alkyl)amino, (C_1-C_4
alkoxy)carbonylamino, (C_1-C_4 alkoxy)carbonyl, hydroxy,
35 C_1-C_4 alkoxy, C_1-C_4 alkylthio or C_1-C_4 alkyl; or

(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, acetamido, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy,

5 C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or trifluoromethyl radicals;

each R₂₁ is independently hydrogen radical or R₂₀;

10 each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or phenyl-C₁-C₂-alkyl or heteroaryl-C₁-C₂-alkyl optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

R₄ is a C₁-C₈ alkyl radical optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy or aryl or heteroaryl

20 optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₄ alkoxy, C₁-C₄ alkylthio, halo, C₁-C₄ alkyl, trifluoromethoxy or trifluoromethyl radicals;

25 R₁₁ is an aryl radical and R₁₂ is a heteroaryl radical, wherein the aryl and heteroaryl radicals are optionally substituted by 1-2 radicals of

(1) R₃₀;

(2) halo or cyano radicals; or

30 (3) -C(O)-NR₃₁R₃₂, -OR₂₉, -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -NR₃₁R₃₂ or -NR₃₃-C(O)-R₂₉ radicals; provided that the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each of R₁₁ and R₁₂ is 0-1;

each R₃₀ is independently

- (1) C₁-C₄ alkyl radical optionally substituted by a phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;
- 5 (2) trifluoromethyl radical; or
- (3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;

each R₂₉ is independently hydrogen radical or R₃₀;

15 each R₃₁ is independently hydrogen, methyl or ethyl radicals; and

each R₃₂ is independently

- (1) hydrogen radicals;
- 20 (2) C₁-C₄ alkyl radical or C₁-C₂ alkyl radical substituted by phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals; or
- 25 (3) phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals.

30 15. The compound of Claim 14 or a pharmaceutically acceptable salt thereof, wherein

R₄ is a C₁-C₄ alkyl radical;

35 R₁₁ is an aryl radical optionally substituted by 1-2 radicals of

- (1) R₃₀;
- (2) halo or cyano radicals; or
- (3) -C(O)-NR₃₁R₃₂, -OR₂₉, -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -NR₃₁R₃₂ or -NR₃₃-C(O)-R₂₉ radicals; and

5

R₁₂ is a heteroaryl radical optionally substituted by 1-2 radicals of

- (1) R₃₀;
- (2) halo or cyano radicals; or
- 10 (3) -C(O)-NR₃₁R₃₂, -OR₂₉, -SR₂₉, -NR₃₁R₃₂ or -NR₃₃-C(O)-R₂₉ radicals;

provided that the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each of R₁₁ and R₁₂ is 0-1;

15

R₃₀ is independently

- (1) C₁-C₄ alkyl radical optionally substituted by a phenyl or heteroaryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;
- (2) trifluoromethyl radical; or
- (3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;

R₂₉ is an aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals; and

R₃₂ is independently

- (1) hydrogen or C₁-C₄ alkyl radical; or

(2) phenyl or heteroaryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals.

5

16. The compound of Claim 15 or a pharmaceutically acceptable salt thereof, wherein

wherein R₁ is -Y or -Z-Y, provided that (1) the total
10 number of aryl, heteroaryl, cycloalkyl and heterocyclyl
radicals in R₁ is 0-1;

15 Z is a C₁-C₄ alkyl radical optionally substituted by 1-2
radicals of amino, di-(C₁-C₂ alkyl)amino, (C₁-C₄
alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂
alkylthio, halo, or aryl or heteroaryl optionally
substituted by 1-2 radicals of hydroxy, C₁-C₂ alkoxy,
C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or
trifluoromethyl radicals;

20

each R₅ is independently hydrogen or C₁-C₄ alkyl
radical;

each R₂₀ is independently

25 (1) C₁-C₈ alkyl radicals optionally substituted by 1-3
radicals of -CO₂R₂₃, amino, C₁-C₄ alkylamino, di-(C₁-C₄
alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-
C₄ alkyl)amino, aminocarbonylamino, hydroxy, C₁-C₄
30 alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄
alkylsulfonyl, halo or C₃-C₆ cycloalkyl, heterocyclyl,
aryl or heteroaryl radicals optionally substituted by 1-
2 radicals of amino, di-(C₁-C₄ alkyl)amino, C₁-C₅
alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄
35 alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-

C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

(2) heterocyclyl radical optionally substituted by 1-2 radicals of (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄

5 alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl; or

(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of (C₁-C₄ alkoxy)carbonyl, amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl

10 or trifluoromethyl radicals;

each R₂₁ is independently hydrogen radical or R₂₀;

each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or

15 phenyl-C₁-C₂-alkyl optionally substituted by 1-2 radicals of hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

R₄ is a methyl or ethyl radical;

20 R₁₁ is an aryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methylthio, methylsulfinyl, methylsulfonyl, aminocarbonyl, methyl or trifluoromethyl

25 radicals; and

R₁₂ is a heteroaryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methyl or trifluoromethyl

30 radicals.

17. The compound of Claim 16 or a pharmaceutically acceptable salt thereof, wherein

Z is C₁-C₄ alkyl radical optionally substituted by 1-2 radicals of amino, t-butoxycarbonylamino, dimethylamino, hydroxy, methoxy, methylthio or halo radicals;

5 Y is a

- (1) hydrogen radical;
- (2) -C(O)-R₂₀, -C(O)-OR₂₁ or -C(O)-NR₅R₂₁ radical;
- (3) -OR₂₁, -SR₂₁, -S(O)-R₂₀, -S(O)₂-R₂₀ or -S(O)₂-NR₅R₂₁ radical; or

10 (4) -NR₅R₂₁, -NR₂₂-C(O)-R₂₁ or -NR₂₂-S(O)₂-R₂₀ radical;

R₅ is a hydrogen radical;

each R₂₀ is independently

15 (1) C₁-C₆ alkyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, methylamino, dimethylamino, t-butoxycarbonylamino, N-((t-butoxy)carbonyl)-N-(methyl)amino, aminocarbonylamino, hydroxy, butoxy, methoxy, butylthio, methylthio, methylsulfinyl, methylsulfonyl, halo or C₅-C₆ cycloalkyl, heterocyclyl, phenyl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, acetamino, hydroxy, methoxy, methylthio, halo, methyl or trifluoromethyl radicals;

20 (2) heterocyclyl radical optionally substituted by 1-2 radicals of t-butoxycarbonyl, hydroxy, or C₁-C₄ alkyl; or

25 (3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, hydroxy, methoxy, methylthio, halo, methyl or trifluoromethyl radicals;

30 each R₂₁ is independently hydrogen radical or R₂₀;

35 each R₂₂ is independently hydrogen or methyl radical;

each R₂₃ is independently hydrogen or C₁-C₄ alkyl radicals;

R₁₁ is an unsubstituted phenyl or naphthyl radical or a
5 phenyl radical substituted by 1-2 radicals of amino,
dimethylamino, acetamido, hydroxy, halo, cyano, methoxy,
methylthio, methylsulfinyl, methylsulfonyl,
aminocarbonyl, methyl or trifluoromethyl radicals; and

10 R₁₂ is a 4-pyridyl, 4-quinolinyl, 4-imidazolyl or 4-pyrimidinyl radical optionally substituted by a radical of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methyl or trifluoromethyl radicals.

15

18. The compound of Claim 17 or a pharmaceutically acceptable salt thereof, wherein

Y is a

20 (1) -C(O)-R₂₀ or -C(O)-NR₅R₂₁ radical;
(2) -OR₂₁, -SR₂₁, -S(O)-R₂₀, -S(O)₂-R₂₀ or -S(O)₂-NR₅R₂₁
radical; or
(3) -NR₅R₂₁, -NR₂₂-C(O)-R₂₁ or -NR₂₂-S(O)₂-R₂₀ radical;

25 each R₂₀ is independently
(1) C₁-C₆ alkyl radicals optionally substituted by 1-3
radicals of -CO₂R₂₃, amino, methylamino, dimethylamino,
t-butoxycarbonylamino, N-((t-butoxy)carbonyl)-N-
(methyl)amino, aminocarbonylamino, hydroxy, butoxy,
30 methoxy, butylthio, methylthio, methylsulfinyl,
methylsulfonyl, halo or C₅-C₆ cycloalkyl, heterocyclyl,
phenyl or heteroaryl radicals optionally substituted by
1-2 radicals of amino, dimethylamino, acetamino,
hydroxy, methoxy, methylthio, halo, methyl or
35 trifluoromethyl radicals;
(2) heterocyclyl radical optionally substituted by t-
butoxycarbonyl; or

(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, hydroxy, methoxy, methylthio, halo, methyl or trifluoromethyl radicals; and

5

each R₂₁ is independently hydrogen radical or R₂₀.

19. The compound of Claim 18 or a pharmaceutically
10 acceptable salt thereof, wherein

Y is a -OR₂₁, -SR₂₁ or -NR₅R₂₁ radical;

each R₂₀ is independently

15 (1) C₁-C₆ alkyl radicals optionally substituted by 1-3 radicals of amino, methylamino, dimethylamino, hydroxy or phenyl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, hydroxy, methoxy, methylthio, halo, methyl or trifluoromethyl
20 radicals;

(2) heterocyclyl radical; or

(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, hydroxy, methoxy, methylthio, halo, methyl or trifluoromethyl
25 radicals;

each R₂₁ is independently hydrogen radical or R₂₀;

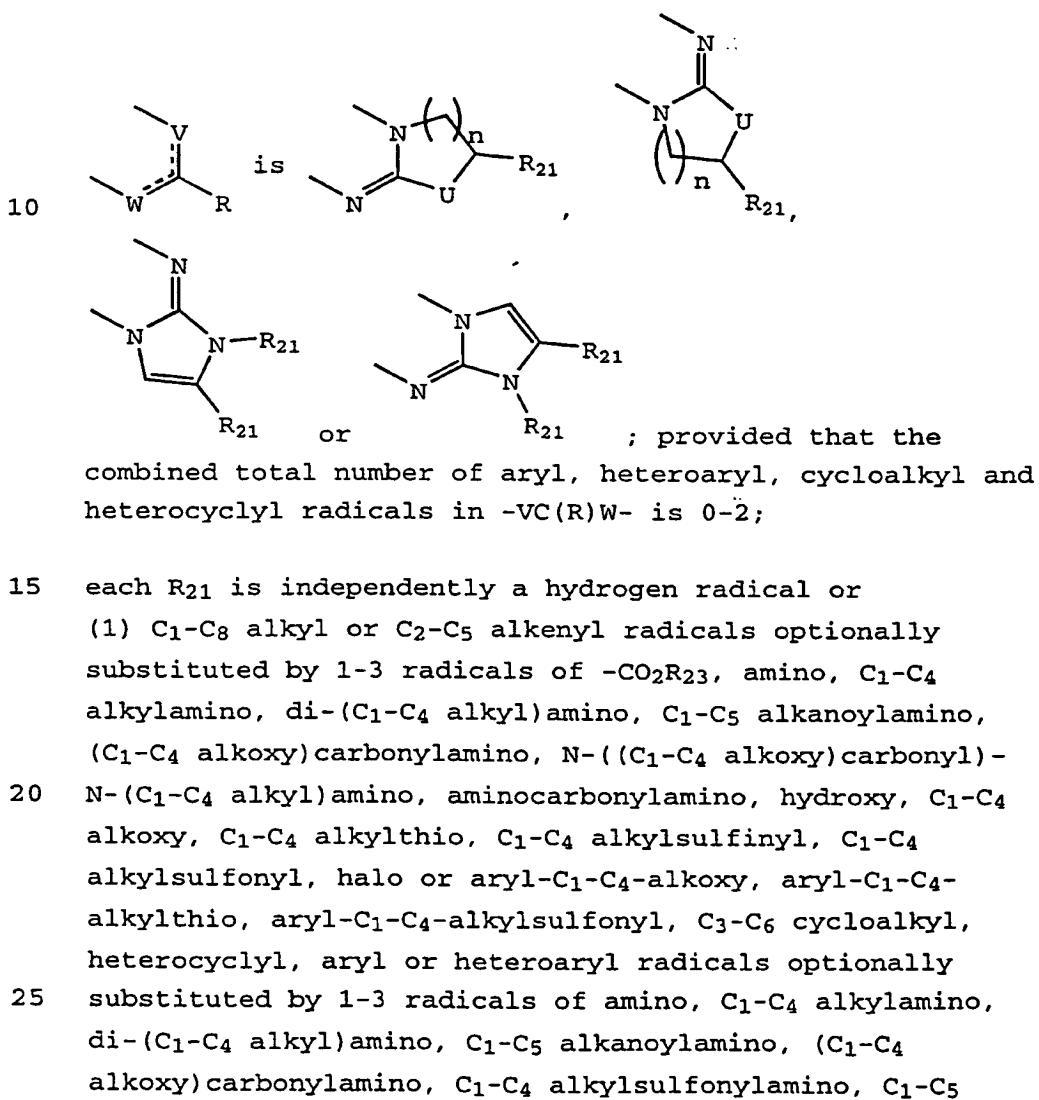
30 R₁₁ is an unsubstituted phenyl radical or a phenyl radical substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methylthio, methylsulfonyl, methyl or trifluoromethyl radicals; and

35 R₁₂ is a 4-pyridyl radical optionally substituted by a radical of amino, dimethylamino, acetamido, hydroxy,

halo, cyano, methoxy, methyl or trifluoromethyl radicals.

5 20. The compound of Claim 3 or a pharmaceutically acceptable salt thereof, wherein

X is O or S;



alkanoyl, (C_1-C_4 alkoxy)carbonyl, hydroxy, C_1-C_4 alkoxy, C_1-C_4 alkylthio, cyano, halo, C_1-C_4 alkyl or C_1-C_2 haloalkyl of 1-3 halo radicals;

(2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, C_1-C_4 alkylamino, di-(C_1-C_4 alkyl)amino, C_1-C_5 alkanoylamino, (C_1-C_4 alkoxy)carbonylamino, (C_1-C_4 alkoxy)carbonyl, hydroxy, C_1-C_4 alkylthio or C_1-C_4 alkyl; or

(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C_1-C_4 alkylamino, di-(C_1-C_4 alkyl)amino, C_1-C_5 alkanoylamino, (C_1-C_4 alkoxy)carbonylamino, C_1-C_4 alkylsulfonylamino, (C_1-C_4 alkoxy)carbonyl, hydroxy, C_1-C_4 alkoxy, C_1-C_4 alkylthio, cyano, halo, azido, C_1-C_4 alkyl or C_1-C_2 haloalkyl of 1-3 halo radicals;

each R_{23} is independently hydrogen or C_1-C_4 alkyl, or phenyl, heteroaryl, phenyl- C_1-C_2 -alkyl or heteroaryl- C_1-C_2 -alkyl optionally substituted by 1-3 radicals of amino, di-(C_1-C_4 alkyl)amino, C_1-C_5 alkanoylamino, (C_1-C_4 alkoxy)carbonylamino, hydroxy, C_1-C_4 alkoxy, C_1-C_4 alkylthio, cyano, halo, C_1-C_4 alkyl or C_1-C_2 haloalkyl of 1-3 halo radicals;

R_{11} and R_{12} are each independently an aryl or heteroaryl radical optionally substituted by 1-2 radicals of

(1) R_{30} ;

(2) halo or cyano radicals;

(3) $-C(O)-R_{30}$, $-C(O)-OR_{29}$, $-C(O)-NR_{31}R_{32}$ or $-C(NR_{31})-$

$NR_{31}R_{32}$ radicals; or

(4) $-OR_{29}$, $-SR_{29}$, $-S(O)-R_{30}$, $-S(O)_2-R_{30}$, $-S(O)_2-NR_{31}R_{32}$, $-NR_{31}R_{32}$, $-NR_{33}-C(O)-R_{29}$ or $-NR_{33}-C(O)-OR_{30}$ radicals;

provided that (1) R_{11} is other than a 4-pyridyl, 4-pyrimidinyl, 4-quinolyl or 6-isoquinoliny radical

optionally substituted by 1-2 substituents; and (2) the

total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each of R₁₁ and R₁₂ is 0-1;

5 each R₃₀ is independently
(1) C₁-C₄ alkyl radical optionally substituted by
(a) amino, C₁-C₄ alkylamino or di-(C₁-C₄-alkyl)amino
radicals; or
(b) hydroxy, C₁-C₄ alkoxy, heterocyclyl, phenyl or
10 heteroaryl radicals optionally substituted by 1-3
radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄
alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄
alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl
15 radicals;
(2) C₁-C₂ haloalkyl of 1-3 halo radical; or
(3) aryl or heteroaryl radicals optionally substituted
by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄
alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄
20 alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl
radicals;

each R₂₉ is independently hydrogen radical or R₃₀;
25 each R₃₁ is independently hydrogen or C₁-C₄ alkyl
radicals; and

each R₃₂ is independently
30 (1) hydrogen radicals;
(2) C₁-C₄ alkyl radical optionally substituted by phenyl
or heteroaryl radical optionally substituted by 1-3
radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄
alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
35 alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl
or trifluoromethyl radicals; or

(3) phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkyl
5 or trifluoromethyl radicals; and

each R₃₃ is independently hydrogen or methyl radical;
and

10 wherein heterocyclyl is a radical of a monocyclic
saturated heterocyclic ring system having 5-6 ring
members, wherein 1-3 ring members are oxygen, sulfur or
nitrogen heteroatoms, which is optionally benzo-fused
and optionally substituted by 1-2 oxo or thioxo,
15 radicals; aryl is a phenyl or naphthyl radical; and
heteroaryl is radical of a monocyclic aromatic
heterocyclic ring system having 5-6 ring members,
wherein 1-3 ring members are oxygen, sulfur or nitrogen
heteroatoms, which is optionally benzo-fused or
20 saturated C₃-C₄-carbocyclic-fused.

21. The compound of Claim 20 or a pharmaceutically
acceptable salt thereof, wherein

25

U is NR₂₁;

each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or
phenyl, heteroaryl, phenyl-C₁-C₂-alkyl or heteroaryl-C₁-
30 C₂-alkyl optionally substituted by 1-3 radicals of
amino, di-(C₁-C₂ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄
alkoxy)carbonylamino, hydroxy, C₁-C₂ alkoxy, C₁-C₂
alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl
radicals;

35

R₁₁ is an aryl radical and R₁₂ is a heteroaryl radical, wherein the aryl and heteroaryl radicals are optionally substituted by 1-2 radicals of

- (1) R₃₀;
- 5 (2) halo or cyano radicals;
- (3) -C(O)-R₃₀, -C(O)-OR₂₉, -C(O)-NR₃₁R₃₂ or -C(NR₃₁)-NR₃₁R₃₂ radicals; or
- (4) -OR₂₉, -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -NR₃₁R₃₂ or -NR₃₃-C(O)-R₂₉ radicals;
- 10 provided that the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each of R₁₁ and R₁₂ is 0-1;

each R₃₀ is independently

- 15 (1) C₁-C₄ alkyl radical optionally substituted by a phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, hydroxy, C₁-C₂ alkoxy, halo, C₁-C₄ alkyl or trifluoromethyl radicals;
- 20 (2) trifluoromethyl radical; or
- (3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, hydroxy, C₁-C₂ alkoxy, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

25 each R₂₉ is independently hydrogen radical or R₃₀; and

each R₃₂ is independently

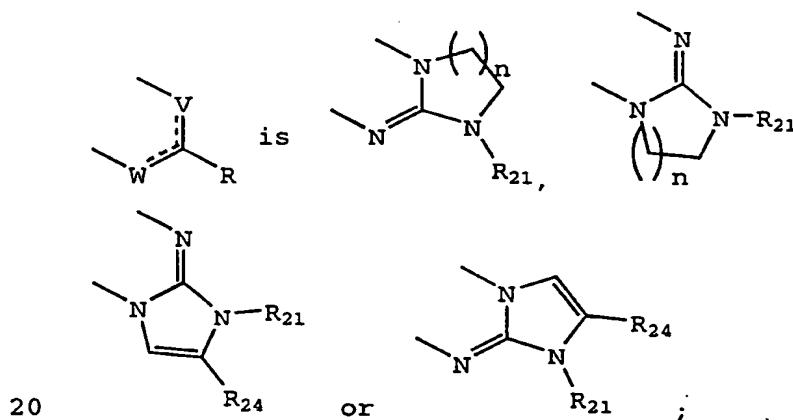
- (1) hydrogen radicals;
- 30 (2) C₁-C₄ alkyl radical or C₁-C₂ alkyl radical substituted by phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, hydroxy, C₁-C₂ alkoxy, C₁-C₄ alkyl or trifluoromethyl radicals; or
- 35 (3) phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino,

acetamido, hydroxy, C₁-C₂ alkoxy, C₁-C₄ alkyl or trifluoromethyl radicals; and

wherein heterocyclyl is a radical of a monocyclic
 5 saturated heterocyclic ring system having 5-6 ring members, wherein 1-2 ring members are oxygen, sulfur or nitrogen heteroatoms, which is optionally benzo-fused and optionally substituted by 1-2 oxo or thioxo radicals; aryl is a phenyl or naphthyl radical; and
 10 heteroaryl is radical of a monocyclic aromatic heterocyclic ring system having 5-6 ring members, wherein 1-2 ring members are oxygen, sulfur or nitrogen heteroatoms, which is optionally benzo-fused.

15

22. The compound of Claim 21 or a pharmaceutically acceptable salt thereof, wherein



each R₂₁ is independently a hydrogen radical or
 (1) C₁-C₈ alkyl or C₂-C₅ alkenyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, C₁-C₄
 25 alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-C₄ alkyl)amino, aminocarbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄

alkylsulfonyl, halo or aryl-C₁-C₄-alkoxy, aryl-C₁-C₄-alkylthio, aryl-C₁-C₄-alkylsulfonyl, C₃-C₆ cycloalkyl, heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, C₁-C₄ alkylamino,
5 di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, C₁-C₅ alkanoyl, (C₁-C₄ alkoxy)carbonyl; hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or C₁-C₂ haloalkyl of 1-3 halo radicals;
10 (2) heterocyclyl radical optionally substituted by 1-2 radicals of amino, di-(C₁-C₄ alkyl)amino, (C₁-C₄ alkoxy)carbonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl; or
15 (3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, acetamido, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or trifluoromethyl radicals;
20 each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or phenyl-C₁-C₂-alkyl or heteroaryl-C₁-C₂-alkyl optionally substituted by 1-3 radicals of amino, di-(C₁-C₂ alkyl)amino, acetamido, (C₁-C₄ alkoxy)carbonylamino,
25 hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;
each R₂₄ is independently a hydrogen or C₁-C₄ alkyl radical;
30 R₁₁ is an aryl radical and R₁₂ is a heteroaryl radical, wherein the aryl and heteroaryl radicals are optionally substituted by 1-2 radicals of
(1) R₃₀;
35 (2) halo or cyano radicals; or

(3) $-C(O)-NR_{31}R_{32}$, $-OR_{29}$, $-SR_{29}$, $-S(O)-R_{30}$, $-S(O)_2-R_{30}$, $-S(O)_2-NR_{31}R_{32}$, $-NR_{31}R_{32}$ or $-NR_{33}-C(O)-R_{29}$ radicals; provided that the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each 5 of R_{11} and R_{12} is 0-1;

each R_{30} is independently

10 (1) C_1-C_4 alkyl radical optionally substituted by a phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;

(2) trifluoromethyl radical; or

15 (3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;

each R_{29} is independently hydrogen radical or R_{30} ;

20 each R_{31} is independently hydrogen, methyl or ethyl radicals; and

each R_{32} is independently

25 (1) hydrogen radicals;

(2) C_1-C_4 alkyl radical or C_1-C_2 alkyl radical substituted by phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals; or

30 (3) phenyl or heteroaryl radical optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals.

35

23. The compound of Claim 22 or a pharmaceutically acceptable salt thereof, wherein

R₁₁ is an aryl radical optionally substituted by 1-2 radicals of

(1) R₃₀;

5 (2) halo or cyano radicals; or

(3) -C(O)-NR₃₁R₃₂, -OR₂₉, -SR₂₉, -S(O)-R₃₀, -S(O)₂-R₃₀, -S(O)₂-NR₃₁R₃₂, -NR₃₁R₃₂ or -NR₃₃-C(O)-R₂₉ radicals; and

R₁₂ is a heteroaryl radical optionally substituted by 1-

10 2 radicals of

(1) R₃₀;

(2) halo or cyano radicals; or

(3) -C(O)-NR₃₁R₃₂, -OR₂₉, -SR₂₉, -NR₃₁R₃₂ or -NR₃₃-C(O)-R₂₉ radicals;

15 provided that the total number of aryl, heteroaryl, cycloalkyl and heterocyclyl radicals substituted on each of R₁₁ and R₁₂ is 0-1;

R₃₀ is independently

20 (1) C₁-C₄ alkyl radical optionally substituted by a phenyl or heteroaryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;

25 (2) trifluoromethyl radical; or

(3) aryl or heteroaryl radicals optionally substituted by 1-3 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals;

30 R₂₉ is an aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, methoxy, methyl or trifluoromethyl radicals; and

35

R₃₂ is independently

- (1) hydrogen or C₁-C₄ alkyl radical; or
- (2) phenyl or heteroaryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, methoxy, methyl or trifluoromethyl radicals.

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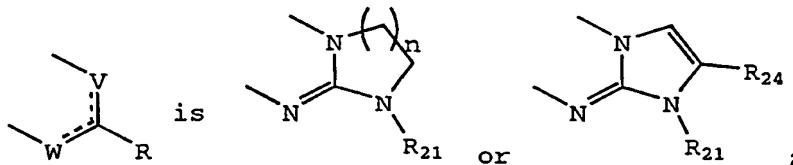
24. The compound of Claim 23 or a pharmaceutically acceptable salt thereof, wherein

- 10 each R₂₁ is independently a hydrogen radical or
 - (1) C₁-C₈ alkyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, N-((C₁-C₄ alkoxy)carbonyl)-N-(C₁-C₄ alkyl)amino, aminocarbonylamino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, halo or C₃-C₆ cycloalkyl, heterocyclyl, aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, di-(C₁-C₄ alkyl)amino, C₁-C₅ alkanoylamino, (C₁-C₄ alkoxy)carbonylamino, C₁-C₄ alkylsulfonylamino, (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;
 - (2) heterocyclyl radical optionally substituted by 1-2 radicals of (C₁-C₄ alkoxy)carbonyl, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio or C₁-C₄ alkyl; or
 - (3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of (C₁-C₄ alkoxy)carbonyl, amino, C₁-C₄ alkylamino, di-(C₁-C₄ alkyl)amino, hydroxy, C₁-C₄ alkoxy, C₁-C₄ alkylthio, cyano, halo, azido, C₁-C₄ alkyl or trifluoromethyl radicals;
- 25 each R₂₃ is independently hydrogen or C₁-C₄ alkyl, or phenyl-C₁-C₂-alkyl optionally substituted by 1-2 radicals of hydroxy, C₁-C₂ alkoxy, C₁-C₂ alkylthio, cyano, halo, C₁-C₄ alkyl or trifluoromethyl radicals;

R₁₁ is an aryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methylthio, methylsulfinyl, 5 methylsulfonyl, aminocarbonyl, methyl or trifluoromethyl radicals; and

R₁₂ is a heteroaryl radical optionally substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, 10 halo, cyano, methoxy, methyl or trifluoromethyl radicals.

25. The compound of Claim 24 or a pharmaceutically acceptable salt thereof, wherein



each R₂₁ is independently a hydrogen radical or
20 (1) C₁-C₆ alkyl radicals optionally substituted by 1-3 radicals of -CO₂R₂₃, amino, methylamino, dimethylamino, t-butoxycarbonylamino, N-((t-butoxy)carbonyl)-N-(methyl)amino, aminocarbonylamino, hydroxy, butoxy, methoxy, butylthio, methylthio, methylsulfinyl, 25 methylsulfonyl, halo or C₅-C₆ cycloalkyl, heterocyclyl, phenyl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, acetamino, hydroxy, methoxy, methylthio, halo, methyl or trifluoromethyl radicals;
30 (2) heterocyclyl radical optionally substituted by t-butoxycarbonyl; or
(3) aryl or heteroaryl radicals optionally substituted by 1-2 radicals of amino, dimethylamino, hydroxy,

methoxy, methylthio, halo, methyl or trifluoromethyl radicals;

each R₂₃ is independently hydrogen or C₁-C₄ alkyl
5 radicals;

R₁₁ is an unsubstituted phenyl or naphthyl radical or a phenyl radical substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, 10 methylthio, methylsulfinyl, methylsulfonyl, aminocarbonyl, methyl or trifluoromethyl radicals; and

R₁₂ is a 4-pyridyl, 4-quinolinyl, 4-imidazolyl or 4-pyrimidinyl radical optionally substituted by a radical 15 of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy, methyl or trifluoromethyl radicals.

26. The compound of Claim 25 or a pharmaceutically acceptable salt thereof, wherein

each R₂₁ is independently a hydrogen radical or
(1) C₁-C₆ alkyl radicals optionally substituted by 1-3
radicals of amino, methylamino, dimethylamino, hydroxy
25 or phenyl or heteroaryl radicals optionally substituted
by 1-2 radicals of amino, dimethylamino, hydroxy,
methoxy, methylthio, halo, methyl or trifluoromethyl
radicals;
(2) heterocyclyl radical; or
30 (3) aryl or heteroaryl radicals optionally substituted
by 1-2 radicals of amino, dimethylamino, hydroxy,
methoxy, methylthio, halo, methyl or trifluoromethyl
radicals;

35 R₁₁ is an unsubstituted phenyl radical or a phenyl radical substituted by 1-2 radicals of amino, dimethylamino, acetamido, hydroxy, halo, cyano, methoxy,

methylthio, methylsulfonyl, methyl or trifluoromethyl radicals; and

R₁₂ is a 4-pyridyl radical optionally substituted by a
5 radical of amino, dimethylamino, acetamido, hydroxy,
halo, cyano, methoxy, methyl or trifluoromethyl
radicals.

10 27. The compound of Claim 1 which is:

2-(2,6-Dichlorobenzyl)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
2-(Butylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
15 2-(Benzylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
5-(4-Fluorophenyl)-3-methyl-((R-1-phenylethyl)amino)-(4-pyridyl)-4(3H)-pyrimidinone,
2-(2-(2-Chlorophenyl)-ethylamino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
20 5-(4-Fluorophenyl)-2-(2-(4-fluorophenyl)-ethylamino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
5-(4-Fluorophenyl)-2-((2-hydroxy-2-phenyl)-ethylamino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
25 5-(4-Fluorophenyl)-3-methyl-2-((3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone,
5-(4-Fluorophenyl)-3-methyl-2-((1-methyl-3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone,
30 5-(4-Fluorophenyl)-3-methyl-2-((R-1-methyl-3-phenylpropyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone,
5-(4-Fluorophenyl)-3-methyl-2-((2-phenylaminoethyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone,
5-(4-Fluorophenyl)-2-((3-imidazolylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
35 5-(4-Fluorophenyl)-3-methyl-6-(4-pyridyl)-2-(3-pyrrolidin-1-yl)-propylamino)-4(3H)-pyrimidinone,
3,6-Diphenyl-4-(4-pyridyl)-2(1H)-pyridone,
6-(4-Methylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone,
40 6-(4-Ethylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone,
6-(2,4-Dimethylphenyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone,

3-Phenyl-4-(4-pyridyl)-6-(2-thienyl)-2(1H)-pyridone,
 6-(2-Furyl)-3-phenyl-4-(4-pyridyl)-2(1H)-pyridone,
 2-(((S)-2-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
 5 2-(((R)-2-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
 2-(((S)-2-N-Ethyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
 10 2-((2-Amino-2-methyl-3-phenylpropyl)amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
 2-((2-Aminomethyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
 2-((3-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
 15 5-(4-Fluorophenyl)-3-methyl-2-((2-methylphenyl)propyl)-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone,
 5-(4-Fluorophenyl)-3-methyl-2-((R,S)-2-amino-3-(2'-fluorophenyl)-propyl-amino)-6-(4-pyridyl)-4(3H)-pyrimidinone,
 20 2-(((S)-2-Acetamido-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
 5-(4-Fluorophenyl)-2-(((S)-2-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
 25 2-(((S)-2-N-n-Butylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
 2-(((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4(3H)-pyrimidinone,
 30 5-(4-Fluorophenyl)-3-methyl-2-((2-methyl-3-phenylpropyl)amino)-6-(4-pyridyl)-4(3H)-pyrimidinone,
 2-(((S)-2-Amino-3-phenylpropyl)-amino)-3-ethyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,
 35 3-Ethyl-5-(4-fluorophenyl)-2-((2-methyl-3-phenylpropyl)amino)-6-(4-pyridyl)-4(3H)-pyrimidinone,
 2-(((S)-2-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-4-(4-pyridyl)-pyrimidine,
 2-((2-(3-trifluoromethylphenyl)phenylmethyl)amino)-3-methyl-5-(4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,
 40 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(4-tolyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,
 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(4-trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-isopropylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-chloro-4-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

5 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3,5-bis(trifluoromethyl)phenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3,4-dichlorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

10 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(1-naphthyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

15 3-Methyl-2-(2(S)-amino-3-phenylpropylamino)-5-(3-trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

3-Methyl-2-(3-phenylpropylamino)-5-(3,5-dichlorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

3-Methyl-2-(3-phenylpropylamino)-5-(4-tolyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

20 3-Methyl-2-(3-phenylpropylamino)-5-(3-trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

3-Methyl-2-(3-phenylpropylamino)-5-(4-methoxyphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

25 3-Methyl-2-(3-phenylpropylamino)-5-(4-trifluoromethylphenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

3-Methyl-2-(2-methyl-3-phenylpropylamino)-5-(3-fluorophenyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

3-Methyl-2-(2-methyl-3-phenylpropylamino)-5-(1-naphthyl)-6-(4-pyridyl)-4(3H)-pyrimidinone,

30 5-(4-Fluorophenyl)-2-(((S)-2-N-glycylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

2-(((S)-2-N-Glycylamino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone,

35 5-(4-Fluorophenyl)-2-(((S)-2-hydroxyacetamido-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

5-(4-Fluorophenyl)-2-(((S)-2-pyrrolidinyl-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

40 2-((S)-3-Benzylpiperazinyl)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

2-((3-Amino-3-(2-fluorophenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

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2-((3-Amino-3-(2-methylphenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

5 2-(((S)-3-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

2-(((R)-3-Amino-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

10 2-(((S)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone,

2-(((R)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone,

15 2-((3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone,

2-((3-Amino-3-(2-methylphenyl)propyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone,

20 2-((3-Amino-3-(2-fluorophenyl)propyl)-amino)-3-methyl-6-(4-pyridyl)5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone,

2-((3-Amino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone,

25 2-((3-Amino-3-(2-fluorophenyl)propyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone,

2-((3-Amino-3-(2-chlorophenyl)propyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone,

2-(((S)-3-Amino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)5-(3,4-dimethylphenyl)-4-(3H)-pyrimidinone,

30 2-(((2R,3R)-3-Amino-2-methyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

2-(((2S,3S)-3-Amino-2-methyl-3-phenylpropyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

35 5-(4-Fluorophenyl)-2-(((S)-3-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

5-(4-Fluorophenyl)-2-(((R)-3-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

40 5-(4-Fluorophenyl)-3-methyl-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-4-(3H)-pyrimidinone,

45 3-Methyl-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-5-(3-trifluoromethylphenyl)-4-(3H)-pyrimidinone,

3-Methyl-5-(3-methylphenyl)-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-4-(3H)-pyrimidinone,

5 3-Methyl-5-(4-methylthiophenyl)-6-(4-pyridyl)-2-((S)-tetrahydroisoquinol-3-ylmethylenamino)-4-(3H)-pyrimidinone,

10 2-(((S)-2-Amino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone,

15 5-(4-Fluorophenyl)-2-((3-hydroxy-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

20 2-(((S)-2-Amino-3-(4-fluorophenyl)propyl)-amino)-5-(4-fluorophenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone,

25 2-(((S)-2-Amino-3-(2-fluorophenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

30 2-(((S)-2-Amino-3-(4-fluorophenyl)propyl)-amino)-5-(4-fluorophenyl)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

35 2-(((S)-2-N-Isopropylamino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone,

40 5-(3-Chlorophenyl)-2-(((S)-2-N-isopropylamino-3-phenylpropyl)-amino)-3-methyl-6-(4-pyridyl)-4-(3H)-pyrimidinone,

45 2-(((S)-2-N,N-Dimethylamino-3-phenylpropyl)-amino)-3-methyl-5-(3-methylphenyl)-6-(4-pyridyl)-4-(3H)-pyrimidinone or
5-(4-Fluorophenyl)-3-methyl-2-(((S)-2-N-methylamino-3-phenylpropyl)-amino)-6-(4-pyridyl)-4-(3H)-pyrimidinone or a pharmaceutically acceptable salt thereof.

28. A pharmaceutical composition comprising a compound of Claims 1 to 27 and a pharmaceutically acceptable carrier.

29. A method of prophylaxis or treatment of inflammation comprising administering an effective amount of a compound of Claims 1 to 27.

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30. A method of prophylaxis or treatment of inflammation comprising administering an effective amount of a composition of Claim 28.

10 31. A method of prophylaxis or treatment of rheumatoid arthritis, Pagets disease, osteoporosis, multiple myeloma, uveitis, acute or chronic myelogenous leukemia, pancreatic β cell destruction, osteoarthritis, rheumatoid spondylitis, gouty arthritis, 15 inflammatory bowel disease, adult respiratory distress syndrome (ARDS), psoriasis, Crohn's disease, allergic rhinitis, ulcerative colitis, anaphylaxis, contact dermatitis, asthma, muscle degeneration, cachexia, Reiter's syndrome, type I diabetes, type II diabetes, 20 bone resorption diseases, graft vs. host reaction, Alzheimer's disease, stroke, myocardial infarction, ischemia reperfusion injury, atherosclerosis, brain trauma, multiple sclerosis, cerebral malaria, sepsis, septic shock, toxic shock syndrome, fever, myalgias due 25 to HIV-1, HIV-2, HIV-3, cytomegalovirus (CMV), influenza, adenovirus, the herpes viruses or herpes zoster infection in a mammal comprising administering an effective amount of a compound of Claims 1-27.

30 32. A method of prophylaxis or treatment of rheumatoid arthritis, Pagets disease, osteoporosis, multiple myeloma, uveitis, acute or chronic myelogenous leukemia, pancreatic β cell destruction, osteoarthritis, rheumatoid spondylitis, gouty arthritis, 35 inflammatory bowel disease, adult respiratory distress syndrome (ARDS), psoriasis, Crohn's disease, allergic rhinitis, ulcerative colitis, anaphylaxis, contact

dermatitis, asthma, muscle degeneration, cachexia,
Reiter's syndrome, type I diabetes, type II diabetes,
bone resorption diseases, graft vs. host reaction,
Alzheimer's disease, stroke, myocardial infarction,
5 ischemia reperfusion injury, atherosclerosis, brain
trauma, multiple sclerosis, cerebral malaria, sepsis,
septic shock, toxic shock syndrome, fever, myalgias due
to HIV-1, HIV-2, HIV-3, cytomegalovirus (CMV),
influenza, adenovirus, the herpes viruses or herpes
10 zoster infection in a mammal comprising administering an
effective amount of a composition of Claim 28.

33. A method of lowering plasma concentrations of
either or both TNF-a and IL-1 comprising administering
15 an effective amount of a compound of Claims 1-27.

34. A method of lowering plasma concentrations of
either or both TNF-a and IL-1 comprising administering
an effective amount of a composition of Claim 28.

20 35. A method of lowering plasma concentrations of
either or both IL-6 and IL-8 comprising administering an
effective amount of a compound of Claims 1-27.

25 36. A method of lowering plasma concentrations of
either or both IL-6 and IL-8 comprising administering an
effective amount of a composition of Claim 28.

30 37. A method of prophylaxis or treatment of
diabetes disease in a mammal comprising administering an
effective amount of a compound according to Claims 1 to
27 to produce a glucagon antagonist effect.

35 38. A method of prophylaxis or treatment of
diabetes disease in a mammal comprising administering an
effective amount of a pharmaceutical composition

according to Claim 28 to produce a glucagon antagonist effect.

39. A method of prophylaxis or treatment of a pain
5 disorder in a mammal comprising administering an effective amount of a compound according to Claims 1 to 27.

40. A method of prophylaxis or treatment of a pain
10 disorder in a mammal comprising administering an effective amount of a pharmaceutical composition according to Claim 28.

41. A method of decreasing prostaglandins
15 production in a mammal comprising administering an effective amount of a compound according to Claims 1 to 27.

42. A method of decreasing prostaglandins
20 production in a mammal comprising administering an effective amount of a pharmaceutical composition according to Claim 28.

43. A method of decreasing cyclooxygenase enzyme
25 activity in a mammal comprising administering an effective amount of a compound according to Claims 1 to 27.

44. The method of Claim 43 wherein the
30 cyclooxygenase enzyme is COX-2.

45. A method of decreasing cyclooxygenase enzyme
activity in a mammal comprising administering an
35 effective amount of a pharmaceutical composition according to Claim 28.

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46. The method of Claim 45 wherein the cyclooxygenase enzyme is COX-2.